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Robust Summaries

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Melting Point

Type Melting Point

Test Substance 1,3-Dioxolane
CAS Number: 646-06-o

Method

- Guideline None
- Test Type Melting Point
- GLP No
- Year unknown

Result

- Melting Point -95 deg C

Remarks Field for
Results Handbook data

Conclusions

Remarks Field The melting point is -95°C

Data Quality

- Reliability Klimisch Code 2. A reliability code of 2 is assigned to data from reference handbooks.

References

1. Lide, D.R. (ed). CRC Handbook of Chemistry and Physics. 72nd ed. Boca Raton, FL: CRC Press, 1991-1992.,p. 3-218

Other

Boiling Point

Type Boiling Point

Test Substance 1,3 -Dioxolane
CAS Number: 646-06-o

Method

- Guideline None
- Test Type Boiling Point
- GLP No
- Year Unknown

Result

- Boiling Point 78 deg C @ 765 mm Hg (1)

Remarks Field for
Results Handbook data

Conclusions

Remarks Field Boiling point is 78 deg C @ 765 mm Hg (1)

Data Quality

- Reliability Klimisch Code 2. A reliability code of 2 is assigned to data from reference handbooks.

References

1. Lide, D.R. (ed). CRC Handbook of Chemistry and Physics. 72nd ed. Boca Raton, FL: CRC Press, 1991-1992. p. 3-218


Other

Vapor Pressure

Type Vapor Pressure

Test Substance 1,3-Dioxolane
CAS Number: 646-06-o

Method

- Guideline None
- Test Type Vapor Pressure
- GLP No
-  Unknown

Result

- Vapor Pressure 70 mm Hg @ 20 Deg C

Remarks Field for
Results Handbook data

Conclusions

Remarks Field Vapor pressure is 70 mm Hg at 20°C

Data Quality

- Reliability **Klimisch** Code 2. A reliability code of 2 is assigned to data from reference handbooks.

References

Sax, N.I. and R.J. Lewis, Sr. (eds.). Hawley's Condensed Chemical Dictionary. 11th ed. New York: Van Nostrand Reinhold Co. p 425 (1987).

Other

Partition Coefficient, Octanol-Water

Type Partition Coefficient, Octanol-Water

Test Substance 1,3 -Dioxolane

Method

- Guideline Not specified
- Test Type Partition Coefficient, Octanol-Water
- GLP No
- Year 1995

Result

- Log $K_{o/w}$ Experimental -0.37'
Calculated by KOWWIN -0.31²

Remarks Field for Results Dioxolane was one of the reference compounds for development of the KOWWIN program (module of EPIWIN). The experimental value is from the literature. The calculated value is the result of the KOWWIN calculation. The experimental value is given priority and the calculated value is supporting.

Conclusions

Remarks Field The log $K_{o/w}$ is approximately -0.35. This material is expected to be relatively water soluble and not bioaccumulate to any significant degree.

Data Quality

- Reliability Klimisch Code 2. A reliability code of 2 is generally assigned to literature values not conducted under OECD guidelines or glps.

References

1. Hansch. C., A. Leo and D. Hoekman. Exploring QSAR. Hydrophobic, Electronic, and Steric Constants. ACS Professional Reference Book. Wshington, DC: American Chemical Society. 1995.
2. KOWWIN v 1.66, Syracuse Research Corporation, Syracuse, NY (April 2000)

Other

Water Solubility

Type Water Solubility

Test Substance 1,3-Dioxolane

Method

- **Guideline** None specified
- **Test Type** Water Solubility
- **GLP** No
- **Year** Unknown

Result

- **Solubility** Soluble in water in all proportions,

Remarks Field for Results Handbook data

Conclusions

Remarks Field Material is soluble in water in all proportions.

Data Quality

- **Reliability** Klimisch Code 2. A reliability code of 2 is assigned to data from reference handbooks.

References

1. Lide, D.R. (ed). CRC Handbook of Chemistry and Physics. 72nd ed. Boca Raton, FL: CRC Press, 1991-1992.,p. 3-218

Other

Photodegradation

Type Photodegradation

Test Substance 1,3-Dioxolane

Method

- **Guideline** Estimated Using version 1.90 of the Atmospheric Oxidation Program for Microsoft Windows (AOPWIN)¹ which estimates the rate constant for the atmospheric, gas-phase reaction between photochemically produced hydroxyl radicals and organic chemicals. The rate constants estimated by the program are then used to calculate atmospheric half-lives for organic compounds based upon average atmospheric concentrations of hydroxyl radical.
- **Test Type** Photodegradation Estimate
- **GLP** No
- **Year** 2000

Results

- **Result** APOWIN estimated OH rate constant 11.2 x 10⁻¹² cm³/molecule-sec

Remarks Field for Results The APOWIN estimate for the reaction rate is based on simple hydrogen abstraction. Similar compounds provide estimates close to measured values for this rate constant. Thus, the method is expected to provide an accurate estimate of the reaction rate constant with hydroxyl radical. Based on the estimated rate constant and using the defaults in APOWIN for atmospheric hydroxyl radical concentration, the estimated half-life is approximately 11.5 hours.

Conclusions

Remarks Field The atmospheric half-life of 1,3-Dioxolane in the atmosphere is estimated to be in the range of 11.5 hours

Data Quality

- **Reliability** Klimisch Code 2. A reliability code of 2 is assigned a result using an accepted method of estimation.

References

1. Syracuse Research Corporation, Syracuse, NY (April 2000)

Other

Water Stability

Type Water Stability

Test Substance 1,3-Dioxolane
CAS Number: 646-06-o
Purity 99.98%

Method

- Guideline OECD 111 (1981)
- Test Type Hydrolysis as a Function of pH
- GLP Yes
- Year 2000

Remarks Field for 0 Duration
Test Conditions

Four days (preliminary test only)

0 Analytical
Method

Direct injection into GC using flame ionization detector.

0 Buffers

Target pH	Buffer System	Measured pH at 50° C
4.0	Potassium biphthalate	3.96
7.0	Potassium phosphate	7.03
9.0	Sodium borate	9.04

0 Vessels

Amber Teflon-lined screw cap vials

0 Replicates

Two at each pH.

0 Temperature

51 ± 0.2° c

0 Additional
Testing

Not conducted since material showed less than ten percent degradation at 50° in four days.

Results

- Nominal Target Concentration 400 mg/l

- Measured Concentrations

PH	Measured Concentration (mg/l)		Percent Degradation At 51° C in 4 days
	Initial	Final	
3.96	484	574	2.1
7.03	345	342	0.9
9.04	529	523	1.2

- Percent Degradation Less than 10% at 50" in four days
- Breakdown Products None

Remarks Field for Results Data were consistent at all pH values, there may have been slightly more degradation at the lower pH.

Conclusions

- Remarks Field
- Dioxolane is stable at pH 4, 7 or 9 for four days at 51° C under the conditions specified in the OECD 111 Guideline.
 - Dioxolane is considered to have a half-life ($t_{1/2}$) at 25 ° C greater than one year at pH 4, 7 and 9.

Data Quality

- Reliability Klimisch Code 1. May be used without restriction.

Reference

1,3-Dioxolane CAS No. 646-06-o: Hydrolysis as a Function of pH. Toxikon Corporation, Laboratory Project ID 00J0009f, October 2000.

Other

The hydrolysis-modeling program found in EPIWIN contains no valid model for estimating hydrolysis of ethers.

- References for supporting studies
- HYDROWIN** modeling program, version 1.67, as found in EPIWIN v 3.05, Syracuse Research Corporation, Syracuse NY (April 2000).

Theoretical Distribution (Fugacity)

Type Theoretical Distribution (Fugacity)

Test Substance 1,3-Dioxolane

Method

- **Guideline** Estimated using the **MacKay** model with standard defaults contained in EPIWIN v 3.05.'
- **Test Type** Level III Fugacity Model
- **GLP** No
- **Year** 2000

Result

- **Distribution**
 - o Air 4.1 %
 - o Water 54 %
 - o Soil 42 %
 - o Sediment 0.1 %

Remarks Field for Results This is the currently accepted model for theoretical distribution estimation.

Conclusions

Remarks Field This material is expected to environmentally distribute primarily in water and soil.

Data Quality

- **Reliability** Klimisch Code 2. A reliability code of 2 is assigned a result using an accepted method of estimation.

References

1. Syracuse Research Corporation, Syracuse, NY (April 2000)

Other

Biodegradation

Type Ready Biodegradation

Test Substance 1,3-Dioxolane
CAS Number: 646-06-0

Method

- Guideline OECD 301D
- Test Type Ready Biodegradation
- GLP Yes
- Year 2000
- Contact Time 35 Days
- Innoculum Municipal Water Treatment Secondary Effluent

Remarks for Test Conditions

- ◇ Innoculum
 - Fresh domestic water treatment secondary effluent collected about one week prior to test
 - One ml filtered effluent per liter final concentration.
 - Not acclimated
- ◇ Test Material Concentration
 - 3.04 mg/l
- ◇ Reference Material
 - Sodium Benzoate at 3 mg/l
 - Reference material showed 78% of THOD
- ◇ Incubation Temperature
 - 22° C in the dark
- ◇ Sampling Frequency
 - 0, 3, 5, 11, 14, 17, 21, 28 and 35 days
 - Duplicate bottles sampled
- ◇ Analytical Method
 - Oxygen consumption
- ◇ Controls and Blanks
 - Innoculum blank without test material
 - Positive control using Sodium Benzoate

Result

- Degradation Percent after time 3.7 % Biodegradation in 35 days
- Result Not Readily Biodegradable
- Kinetics Not applicable
- Breakdown Products None determined

Remarks Field for Results Not readily biodegradable

Conclusions

Remarks Field Not readily biodegradable under these conditions

Data Quality

- Reliability Klimisch Code 1. Recent glp study meeting current OECD guideline, may be used without restriction

References

1,3-Dioxolane (CAS 646-06-o): Ready Biodegradability: Closed Bottle Test. Toxikon Corporation September 2000, submitted to Ticona and Ferro Corporations

Other

This study is supported by an earlier study, sponsored by Celanese, in which Dioxolane was tested for biodegradation using a municipal secondary effluent and measuring oxygen consumption with a manometric respirometer. After 15 days Dioxolane showed 5.2 % of the THOD. ¹

This result is supported by the BIOWIN **V4.0** model found in EPIWIN. Two of the three models included predict that Dioxolane will not rapidly **biodegrade**.²

References for Supporting Studies

1. Report to Celanese Chemical Company Inc. on Toxicology and Fate of Selected Industrial Chemicals in Aquatic Ecosystems. J.R. Walton and E.M. Davis, University of Texas at Houston. December 1980.
2. EPIWIN v 3.05, Syracuse Research Corporation, Syracuse NY (April 2000).

Acute Toxicity to Fish

Type Acute Toxicity to Fish

Test Substance **1,3-Dioxolane**
CAS Number: 646-06-o
Purity 99.98%

Method

- Guideline OECD 203 (1992)
- Test Type Acute Toxicity to Fish
- GLP Yes
- Year 2000
- Species/Strain Bluegill sunfish (*Lepomis macrochirus*)
- Analytical Monitoring At the beginning and end of each 24-hour renewal period
- Exposure Period (unit) 96 hours
- Test Details Static renewal, renewal every 24 hour
- Statistical Methods None applied to biology due to lack mortality

Remarks Field for Test Conditions	0 Fish Size and Age	<i>Lepomis macrochirus</i> juveniles, 29 ± 1.9 mm average standard length and 0.57 ± 0.11 g average wet weight.
	0 Test conditions	As specified in OECD 203
	0 Solvent	None, test material water soluble
	0 Dilution Water Source and Chemistry	Filtered city of Jupiter (Florida) freshwater with an initial hardness and alkalinity of 66 and 28 mg/l as CaCO ₃ , respectively and an initial dilution water temperature range of 23.1 to 23.2 °C. Final hardness and alkalinity on day 4 were 74 and 30 mg/l as CaCO ₃ , respectively.

0 Stock and Test Solutions	A stock solution with target concentration of 10,000 mg/kg was prepared by weighing the test material and dissolving it in dilution water. This stock was diluted to prepare the test solutions each day. Concentration of Dioxolane in both the stock and the test dilutions were determined by gc analysis using a glp validated method.
0 Vessels and Lighting	The test chambers were 3.5-gallon glass jars (22-cm diameter x 30-cm height) containing 9.0 L of dilution water and providing a water depth of approximately 25 cm. All test chambers were covered throughout the exposure period to reduce evaporation or contamination. The test chambers were positioned in a water bath set to maintain $23 \pm 2^{\circ}\text{C}$ under fluorescent lighting regulated to a photoperiod of 16 hours light and 8 hours darkness. The light intensity ranged between 11.9 and 17.1 microMols per square meter per second
0 Fish per Vessel and Croup	There were 10 fish in each vessel, 30 fish (3 vessels) per treatment and control
0 Dose Selection	Dose selection of 100 mg/l was based on a range- finding study under static conditions with concentrations as high as 1000 mg/l showing no mortality.
0 Water Chemistry in Control and in One Concentration Where Effects Were Observed	Dissolved oxygen averaged about 8.5 mg/l at the beginning of each renewal period and was found to be between 5.9 and 6.9 mg/l at the end of each 24- hour period. Only one measurement of 48 total was below 6.0 mg/l . The initial pH varied from 7.7 to 8.0 at the beginning of each 24-hour renewal period and ranged from 6.9 to 7.7 at the end of each period (only one measurement of 48 total was below 7.0). The pH and DO were similar in the control and treatment group .
0 Renewal	Solutions were renewed after 24 hours. The test material was known to be volatile.
0 Exposure period and Observations	Four days with daily observations

0 Temperature range 22.0 to 23.2° C as measured at the beginning and end of each 24-hour renewal. The diurnal range of the water bath temperature was continuously monitored using a minimum/maximum thermometer and recorded daily and ranged from 21.6 to 23.4° C over the testing period.

0 Analytical Results

Day	Measured Concentration (mg/l) [†]	
	Initial	24 hours
1	108	93.8
2	102	91.8
3	105	72.1
4	99.6	90.5
	Mean Concentrations* over all measurements 95.4 mg/l	
	Test material below limit of detection in controls	
	† Composite sample from all three vessels	

* Arithmetic mean

Results

- Nominal Concentrations 0,100 mg/l
- Measured Concentrations 95.4 mg/l (mean over 96 hours of test)
- Units mg./l.
- LC₅₀ >95.4 mg/l at 24, 48, 72 and 96 hours.
- LC₀ 95.4 mg/l at 24, 48, 72 and 96 hours
- NOEC 95.4 mg/l at 24, 48, 72 and 96 hours

0 Mortality No fish died during the exposure period

0 Sub-lethal Effects No sub-lethal effects of the test substance were observed.

Conclusions

Remarks Field The 96-hour LC_{50} for the limit test was **>95.4 mg/l** (based on measured concentrations). The no-observed-effect-concentration (**NOEC**) for the limit test was **95.4 mg/l**.

Data Quality

- Reliability Klimisch Code 1. May be used without restriction.

Reference **1,3-Dioxolane** CAS No. 646-06-o: Acute Toxicity to Bluegill, *Lepomis macrochirus*, Under Static-Renewal Test Conditions. Toxikon Corporation, Laboratory Project ID **00J0009g**, October 2000.

Other

- ◇ This study is supported by an earlier study, sponsored by Celanese, in which Sheepshead minnows (*Cyprinodon variegatus*, 5 per group) were exposed to Dioxolane at concentrations of **7500, 11000, 13000, 15000** and **25000 mg/l**. In this study the 48 hour- LC_{50} was reported to be **12000 mg/l**, and the 96-hour LC_0 was **7500 mg/l**, the 96 hour- LC_{50} was reported to be **10000 mg/l**, and the 96-hour LC_0 was **7500 mg/l**. In this study 5 fish per concentration were exposed and a clear dose-response was established with a 24-hour mortality of **5/5** at **25000 mg/l**. Based on the volatility of Dioxolane, the actual value for the EC_{50} in this study is likely somewhat than reported. Nevertheless, this study supports the low order of toxicity found for Dioxolane toward bluegill sunfish 1
- ◇ The EPA ECOSAR Modeling Program found in EPIWIN, estimates the 96-hour LC_{50} for **fish** to be **8150 mg/l**.²

- References for supporting studies
1. Report to Celanese Chemical Company Inc. on Toxicology and Fate of Selected Industrial Chemicals in Aquatic Ecosystems. J.R. Walton and E.M. Davis, University of Texas at Houston. December 1980.
 2. ECOSAR modeling program, version **0.99f**, as found in **EPIWIN v 3.05**, Syracuse Research Corporation, Syracuse NY (April 2000).

Acute Toxicity to Aquatic Invertebrates

Type Acute Toxicity to Aquatic Invertebrates

Test Substance **1,3-Dioxolane**
CAS Number: 646-06-o
Purity 99.98%

Method

- Guideline OECD 202 (1984)
- Test Type Daphnia, acute immobilization
- GLP Yes
- Year 2000
- Analytical Procedures Determination of test material concentration at **0, 24** and 48 hours
- Species/Strain *Daphnia magna*
- Test Details Static renewal
- Statistical Methods None necessary due to lack of greater than 50% mortality

Remarks Field for Test Conditions	0 Age at study initiation	Less than 24 hours
	0 Test conditions	As specified in OECD 202.
	0 Solvent	Moderately hard fresh water
	0 Vessel	Crystallizing dishes 10 cm diameter by 5 cm deep. Vessels were kept covered with glass plate during test to reduce loss of test material. Dishes contained 200 ml of solution
	0 Daphnids per group	Twenty
	0 Daphnids per vessel	Five
	0 Exposure period	48 hours
	0 Temperature range	19.2 to 20.0° C
	0 Observation times	24 and 48 hours
	0 Solution pH range	7.6 to 7.8 at all concentrations at both 0 and 48 hours

0 **Dissolved oxygen** Above 8.8 mg/l for all solutions at 0 and 48 hours

0 Renewal Solutions were renewed after 24 hours

0 Analytical Results

Target Conc	Measured Concentration (mg/l)				
	Initial	24 hours	24 hours Renewed	48 Hours	Mean Conc*
250	244	179	240	190	213
500	485	315	495	348	411
1000	982	546	990	570	772

* Arithmetic mean

0 Water Parameters

- Hardness of 68 mg/l as calcium carbonate
- Alkalinity of 26 mg/l as calcium carbonate
- Specific conductivity of 527 microSiemens

Results

- Nominal Concentrations 0, 250, 500 and 1000 mg/l
- Measured Concentrations 213, 411 and 772 mg/l (mean over 48 hours of test)
- Units mg./l.
- EC₅₀ >764 at 24 hours
> 772 at 48 hours
- EC₁₁
 - 24 hours: 764 mg/l
 - 48 hours: Not determined

Remarks Field for 0 Immobilization Results

Concentration	Cumulative Number Immobilized	
	24 hours	48 hours
0	0	0
213	0	8/20
411	0	6/20
772	0	9/20

Conclusions

Remarks Field

- The EC_{50} (24 hour) was **>764 mg/l** and the EC_0 (24 hour) was **764 mg/l** based on the mean measured concentrations. The EC_{50} (48 hour) was **>772 mg/l** and the EC_{01} was not found based on the mean measured concentrations
- The volatility of the test substance makes it difficult to accurately determine the exposure concentrations. Based on the lack of mortality and the lack of a typical dose-response curve, it appears that the stress of the renewal conditions may have contributed to the immobilization. It is clear that the test material has a low order of toxicity to daphnids. Because of this result and the difficulty in maintaining concentration levels, further studies were not conducted.
- A preliminary range-finding test was conducted as part of this study. In this test, ten daphnids were exposed for 48 hours to nominal concentrations of 0, 0.1, 1.0, 10.0, 100 or 1000 **mg/l** Dioxolane in water. Under these conditions, no immobilization was observed.

Data Quality

- Reliability **Klimisch Code 1.** May be used without restriction.

Reference

1,3-Dioxolane CAS No. 646-06-o: Acute Toxicity to the Water Flea, *Daphnia magna*, Under Static-Renewal Test Conditions. Toxikon Corporation, Laboratory Project ID **00J0009c** October 2000.

Other

- This study is supported by an earlier study, sponsored by Celanese, in which Dioxolane was tested at **1000, 5000, 6500, 8000, 9000, 10000** and **12500 mg/l**. In this study the 24-hour EC_{50} was reported to be **7650 mg/l** and the **48-hour** EC_{50} was reported to be **6950 mg/l**.¹ In this study 20 daphnids were exposed and a clear dose-response relationship was obtained. Based on the volatility of Dioxolane, the actual value for the EC_{50} in this study is likely lower than reported. Nevertheless, this study supports the low order of toxicity found for Dioxolane toward daphnids.
- The EPA ECOSAR Modeling Program found in EPIWIN, estimates the **48-hour** LC_{50} for daphnia to be **7400 mg/l**.²

References for supporting studies

1. Report to Celanese Chemical Company Inc. on Toxicology and Fate of Selected Industrial Chemicals in Aquatic Ecosystems. J.R. Walton and E.M. Davis, University of Texas at Houston. December 1980.
2. ECOSAR modeling program, version **0.99f**, as found in EPIWIN v 3.05, Syracuse Research Corporation, Syracuse NY (April 2000).

Toxicity to Aquatic Plants

Type	Toxicity to Aquatic Plants
None	1,3-Dioxolane CAS Number: 646-06-0 Purity 99.98%
Method	
• Guideline	OECD 20 1
• Test Type	Algae Growth Inhibition
• GLP	Yes
• Year	2000
• Species/Strain	<i>Selenastrum cupricornutum</i> The culture originated from an inoculum received from the Carolina Biological Supply Company (Burlington, NC) and has been maintained in the laboratory since December 3, 1999.
• Element Basis	Number of cells per ml. And area under the growth curve
• Exposure Period	72 hours
• Analytical Monitoring	Yes
• Statistical Methods	<p>0 EC₅₀ values were calculated based on both biomass growth (comparison of area under the growth curves), the E_bC₅₀, and on the average specific growth rate, the E_rC₅₀. EC₅₀ values and their 95 percent confidence limits were estimated by a computer program (U.S. EPA, 1994) for calculating EC values by probit analysis.</p> <p>0 In addition to the EC₅₀ values, a no-observed-effect concentration (NOEC) was calculated by analysis of variance (ANOVA) with statistical differences between cell density means determined by Dunnett's procedure (U.S.EPA, 1988). Statistical differences were determined at a probability level of 0.05.</p> <p>0 Inhibition calculations are based upon a comparison of the areas under the growth curves and are reported using the symbol E_bC₅₀. The 24, 48 and 72-hour E_bC₅₀ values and their 95 percent confidence limits were calculated.</p>

Remarks Field for
Test Conditions

- 0 Test Temperature Range The temperature ranged from 24.4 to 26.8° C.
- 0 Growth Medium Chemistry

The base water for the test medium was deionized water. The base water was enhanced with reagent-grade nutrients as described in ASTM (1994). The **pH** of the test medium was adjusted to 7.5 ± 0.1 prior to the addition of the test substance.

[American Society for Testing and Materials (ASTM). 1990. Standard Guide for Conducting Static **96-Hour** Toxicity Tests with Microalgae. ASTM Designation E1218-90.]
- 0 Dilution Water Source Deionized water **from** the Town of Jupiter Florida, supplemented as above.
- 0 Exposure Vessel Sterile **250-mL** glass Erlenmeyer flasks covered with gas exchange caps containing 100 ml of algal medium.
- 0 Stock Solutions Prepared

Approximately 1 .0 182 g of the chemical was brought to volume in a 100 ml volumetric flask with deionized water to prepare a stock concentration of 10,200 **mg/l**. The following amounts of stock (**1.9, 3.75, 7.5, 15** and 30 ml) were used to make the test concentrations by mixing with **298.1, 296, 292.5, 285,** and 270 ml of freshwater algal media individually.
- ◊ Light Level and Quality Lighting was continuous fluorescent lighting and intensity was measured daily at the surface of the test solutions during the 72-hour exposure period and ranged from 84 to 138 $\mu\text{E}/\text{m}^2/\text{s}$ as measured by a LI-COR, Inc. Model LI-189 light meter equipped with a 2x quantum sensor.
- 0 Test Design
 - Replicates: three replicates for each test concentration. Six replicates were used for the dilution water control.
 - Concentrations were determined by gc using a glp validated method.
 - o Target: 0, 62.5, 125, 250, 500 and 1000 **mg/l**
 - o Mean measured Control (**<3 1 .0**) , 36.9, 81.0, 163, 280 and 877 **mg/l**.

0 Analytical
Determination of Test
Material Concentrations

Nominal Conc	Measured Concentrations mg/l			
	Day- 1	Day - 3	Mean	Percent nominal
Control	ND	ND	---	---
62.5	60.6	13.1	36.9	59
125	124	37.9	81	64.8
250	262	64.7	163	65.2
500	520	39.3	280	56
1000	1027	726	877	87.7

0 Method of calculating
mean

Arithmetic based on composite samples of each replicate for each concentration at study initiation and study termination

0 Exposure period

72 hours

0 Cell Counts

Algal growth was measured by direct cell count using a 0.1mm deep hemacytometer under a compound microscope. Algal counts were conducted on day one and approximately every 24 hours thereafter. Morphological observations were also conducted on the test treatment using a compound microscope to detect abnormal cell morphology and coloration as compared to the control replicates.

Results

- Nominal Concentrations 0, 62.5, 125, 250, 500 and 1000 mg/l
- Measured Concentrations (<31.0) , 36.9, 81.0, 163,280 and 877 mg/l
- Units mg./l.
- EC₅₀ The E_bC₅₀ and E_rC₅₀ (0-72 hours) were >877 mg /l.
- NOEC 877 mg/l (72-hour)

Remarks Field for 0 Biological
Results Observations

After 72 hours of exposure to **1,3-Dioxolane**, the percentage cell growth inhibition (based on area under the growth curve) compared to the control was 19% at the mean measured concentration of **877 mg/l**. The growth curves of both the control and the test solution exhibited a pattern of exponential growth during the **72-hour** growth period. Observations of cell morphology detected no changes in exposed cells as compared to cells in the control media. There was no significant statistical difference between the algal growth of the control and the test solutions

- 0 Daily Cell Counts From Each Replicate These are presented to substantiate that there was no unusual variation between replicates associated with the possible selective volatilization of the test material from individual flasks.

Measured Conc (mg/l)	Cell Numbers (x 10 ⁴)/ml			
	Replicate	24 hrs	48 hrs	72 hrs
Control	A	1.7	25	407
	B	2.1	36	378
	C	1.1	33	280
	D	0.9	43	358
	E	1.8	28	224
	F	1.9	26	289
36.9	A	1.3	13	318
	B	1.3	23	284
	C	2.2	16	218
81	A	1.9	33	329
	B	1.6	20	298
	C	0.9	48	333
163	A	0.4	30	304
	B	0.9	30	336
	C	1.8	56	189
280	A	1.7	29	324
	B	2.0	24	278
	C	1.3	40	291
877	A	2.1	33	229
	B	3.3	37	287
	C	2.1	34	211

0 Mean Cell Density at Each Concentration at Each Time Point

Measured Conc (mg/l)	Mean Cell Numbers (x 10 ⁴)/ml (s.d.)		
	24 hours	48 hours	72 hours
Control	1.6 (0.475)	32 (6.91)	323 (69.4)
36.9	1.6 (0.520)	17 (5.13)	273 (50.8)
81	1.5 (0.513)	34 (14.0)	320 (19.2)
163	1.6 (0.709)	39 (15.0)	276 (77.3)
280	1.7 (0.351)	31 (8.18)	298 (23.7)
877	2.5 (0.693)	35 (2.08)	242 (39.7)

0 Percent Inhibition

Measured Conc (mg/l)	Percent Inhibition		
	0-24hrs	24-48 hrs	48-72 hrs
36.9	0	-47	-21
81	-17	6	0
163	-100	18	-9
280	17	-2	-7
877	150	15	-19

Conclusions

Remarks Field

The E_bC_{50} and E_rC_{50} (0-72 hours) were >877 mg /l (based on measured concentrations). The 72-hour no-observable-effect concentration (NOEC) was 877 mg/l.

The test material was somewhat volatile; however, sufficient Dioxolane remained in the culture flasks (especially at the highest concentration tested) to provide a valid estimate of the growth inhibition potential of the test material to green algae.

Data Quality

- Reliability

Klimisch Code 1. Reliable without restriction. Study was conducted in accord with current OECD guideline under glp conditions. Analytical measurements verified exposure concentrations.

References

1,3-Dioxolane: Toxicity to The Freshwater Green Alga, *Selenastrum cupricornutum*, Under Static Test Conditions. Toxikon Laboratories, Jupiter FL, Project ID 00J0009b, 27 September 2000, submitted to and sponsored by Ticona Corporation and Ferro Corporation.

Other

This study is supported by an earlier study, sponsored by Celanese, in which Trioxane was tested for growth inhibition of *Selenastrum capricornutum*. In this study, algae growth was measured out to 14 days of exposure at levels of 1000, 5000 or 10000 mg/l with counts recorded on days 3, 6, 10 and 14. Significant inhibition was seen only at 5000 mg/l and above 1000 mg/l was determined to be the NOEC. Graphically, the 96-hour EC₅₀ can be determined to be in the range of 4000 mg/l; however, loss of test material may affect this estimate'

The EPA ECOSAR Modeling Program found in EPIWIN, estimates the 96-hour EC₅₀ for green algae to be 4075 mg/l.²

References for supporting studies

1. Report to Celanese Chemical Company Inc. on Toxicology and Fate of Selected Industrial Chemicals in Aquatic Ecosystems. J.R. Walton and E.M. Davis, University of Texas at Houston. December 1980.
2. ECOSAR modeling program, version 0.99f, as found in EPIWIN v 3.05, Syracuse Research Corporation, Syracuse NY (April 2000).

Acute Oral Toxicity

Type Acute Oral Toxicity

Test Substance **1,3-Dioxolane**
CAS Number: 646-06-0
Commercial Grade

Method

- **Guideline** Based on methods described in 16 CFR 1500.3 (c) Consumer Products Safety Commission: Federal Hazardous Substance Act
- **GLP** No
- **Year** 1980
- **Species** Rat
- **Strain** Albino Rats (Sprague-Dawley CD@)
- **Route of administration** Oral Gavage
- **Doses** 2500, 3500, 5000, 7100, 10000 mg/kg
- **Sex** Males and Females
- **Number of Animals/group** Five
- **Vehicle** None, administered neat

Remarks Field for Test Conditions	0	Age at Study Initiation	Unknown, Young Adults At study initiation males weighed 265 to 289 grams and females weighed 233 to 245 grams
	0	Volume administered	1.3 to 9.2 ml/kg body weigh
	0	Post-dose observation period	14 Days

Results

- LD_{50} LD-50 = 5.2 g/kg with 95% confidence limits of 4.3 to 6.1 g/kg

- Number of deaths at each dose level

Dose Level mg/kg	Mortality, males	Mortality, females
2,500	0/5	0/5
3,500	2/5	0/5
5,000	1/5	2/5
7,100	5/5	5/5
10,000	5/5	5/5

Remarks Field for
Results

0 Time of
death

Dose mg/kg	Males (d= day)	Females
3500	d-1, d-1	none
5000	4 hours	d-1, d-1
7100	2 hours, d-1, d-1, d-2, d-2	1 hour, 4 hours, d- 1, d-1, d-1
10,000	2 hours, 4 hours, 6 hours, 6 hours, d-1	1 hour, 2 hours, 2- hours, 4 hours, 4- hours

0 Clinical Signs

Dose mg/kg	Clinical Signs
2,500	<ul style="list-style-type: none"> Ataxia, 1 female from 2 to 4 hours after dosing. Respiratory rate decrease, 3 animals, 2 to 4 hours Motor activity decrease, all animals 1 to 4 hours.
3,500	<ul style="list-style-type: none"> Ataxia, 2 females 2 or 4 hours after dosing. Fine tremors, 1 male, 4-hours after dosing only Respiratory rate decrease, 5 animals, 2 to 4 hours Motor activity decrease, 9/10 animals starting at 1 hour, decreasing to 2 animals at 24 hours. Piloerection, 3 animals, 2 hours to day-2 Prostration, 3 animals, from 2 to 4 hours.
5,000	<ul style="list-style-type: none"> Ataxia, 1 male, only at 2 4 hours after dosing. Fine tremors, 1 female 1- 4 hours after dosing Respiratory rate decrease, 5 animals, 2 to 4 hours Motor activity decrease, 9 animals starting at 1 hour, decreasing to 4 animals at 24 hours. Prostration, 5 animals total, various times from 1 to 24 hours. Hypothermia, 2 animals at 4 or 24 hours after dosing
7,100	<ul style="list-style-type: none"> Respiratory rate decrease, 8 animals total, various 1 to 24 hours after dosing Motor activity decrease, 1 animal 1-2 hours after dosing. Prostration, 8 animals total, various times from 1 to 24 hours. Hypothermia, 4 animals at 4 or 24 hours after dosing
10,000	<ul style="list-style-type: none"> Respiratory rate decrease, 6 animals total, various 1 to 4 hours after dosing Labored breathing, 2 animals at 1 hour after dosing only Prostration, 9 animals total, various times from 1 to 4 hours. Hypothermia, 1 animal at 2 hours after dosing

- 0 Necropsy Findings Necropsy only conducted on animals dying less than 24 hours after dosing and limited to determining if gavage was the cause of death.
- 0 Target Organs No potential target organs were identified in report, CNS may have been affected by solvent narcosis.
- 0 Sex Differences Effects on males and females were similar.

Conclusions

Remarks Field

The oral **LD₅₀** in the rat was determined to be 5,200 **mg/kg**. Solvent induced narcosis may have been the cause of deaths occurring on day-1.

Study documentation is good. Results are consistent with other data for the material, including literature studies.

Data Quality

• Reliability

Klimisch Code 2. **Study** design, conduct and reporting are considered reliable to address the test endpoint although not conducted in accord with glp standards.

References

Acute Oral Toxicity Study in Rats, **Bio/dynamics** Inc., Project No. 6004-79 (06/05/80), Sponsored by The Celanese Corporation, New York, NY.

Other

This study is supported by a study in the literature that reports the acute oral **LD₅₀** in the rat to be 5,800 **mg/kg**¹. This **study** is also supported by a limit test, conducted by Hoechst, that reported the **LD₅₀** to be greater than 2000 **mg/kg**²

References for supporting studies



1. Czajkowska, T, Krysiak, B and **Popińska**, E. Experimental studies of toxic effects of **1,3,5-trioxane** and **1,3-dioxolane**. I. Acute toxic effect. Med Pr; Vol 38, 1987, **P184-90**.
2. **1,3-Dioxolan** - Profung der akuten **oralen** Toxizitäten der Wistar Ratte. Hoechst AG Pharma Research Toxicology Report No. 87.144 1, September 1987.

Acute Inhalation Toxicity

Type Acute Inhalation Toxicity

Test Substance **1,3-Dioxolane**
CAS Number: 646-06-0
Commercial Grade

Method

- Guideline None specified
- GLP No
-  1980
- Species Rat
- Strain Charles River CD@
- Route of administration Whole-body inhalation as vapor
- Doses 0, 37.9, 60.6, 67.9, 88.4 and 201.9 mg/l (nominal concentrations)
-  Males and Females
- Exposure Period Four hours
- Number of Animals/group Five animals of each sex per dose level
- Vehicle Air

Remarks	Field for	0	Age at Study Initiation	Unknown, Young Adults At study initiation males weighed 204 to 295 grams and females weighed 211 to 273 grams
	Test Conditions	0	Doses	Initial dose was 201.9 mg/l (nominal) which caused 100% mortality. Second group was 37.9 mg/l, which caused 0% mortality. Additional groups were added until enough data were accumulated to calculate an accurate LC ₅₀ .
		0	Post-dose observation period	14 Days

Results

- LC₅₀ 68.4 mg/l (95% confidence interval of 61.0 – 766. mg/l)

- Number of deaths at each dose level

Concentration (mg/l)	Number of deaths		
	Males	Females	Total
0	0	0	0
37.9	0	0	0
60.6	0	1	1
67.9	4	1	5
88.4	4	5	9
201.9	5	5	10

Remarks Field for Results

0 Time of death

Conc. (mg/l)	Males and Females (d=day, h=hour)
60.6	6h
67.9	5h, 8h, dl, dl, dl
88.4	4h, 4h, 5h, 5h, 6h, 7h, dl, dl, dl
201.9	All ten within 2 hours

Clinical Signs

37.9 mg/l

Sign	Onset	Duration	# Animals
Lacrimation	30 min	1 day	Most
Labored breathing	2 hours	3 hours	Most
Prostrate	3 hours	3 hours	All
No response to auditory stimuli	3 hours	1 hour	Most
Loss of muscle tone coordination	45 min	< 24 hours	Most

Clinical Signs

60.6

Sign	Onset	Duration	# Animals
Lacrimation	30 min	2 days	Most
Labored breathing	2 hours	6 hours	Some
Rapid breathing	3 hours	3 hours	All
Reduced activity	30 min	< 24 hours	All
Prostrate	2 hours	3 hours	All
No response to auditory stimuli	2 hours	3 hours	All
Loss of muscle tone coordination	1 hour	2 hours	Most

Clinical Signs 67.9 mg/l	Sign	Onset	Duration	# Animals
	Lacrimation	15min	2days	Most
	Labored breathing	2 hours	6 hours	Half
	Rapid breathing	3 hours	<24 hours	All
	Reduced activity	15 min	< 24 hours	All
	Prostrate	2 hours	3 hours	All
	No response to auditory stimuli	45 min	3 hours	All
	Loss of muscle tone coordination	45 min	30 min	Most

Clinical Signs 88.4 mg/l	Sign	Onset	Duration	# Animals
	Lacrimation	15 min	< 24 hours	Most
	Labored breathing	4 hours	< 24 hours	Many
	Rapid breathing	2 hours	3 hours	All
	Reduced activity	15 min	< 24 hours	All
	Prostrate	2 hours	4 hours	All
	No response to auditory stimuli	60 min	4 hours	All
	Loss of muscle tone coordination	45 min	2 hours	Most

Clinical Signs 201.9 mg/l	Sign	Onset	Duration	# Animals
	Lacrimation	15 min	< 2 hours*	All
	Labored breathing	45 min	< 2 hours*	All
	Salivation	14 min	< 2 hours*	All
	Reduced activity	15 min	< 2 hours*	All
	No response to auditory stimuli	45 min	< 2 hours*	All
	Dead	< 2 hours		All

* Animals died after 2 hours of exposure

<u>Body Weights</u> Group	Day Mean Body Weights (grams)						
		<u>0</u>	<u>1</u>	<u>2</u>	<u>4</u>	<u>7</u>	<u>14</u>
Control	Males	282	285	285	297	314	345
	Females	227	225	228	231	236	244
37.9 mg/l	Males	285	264	263	276	290	318
	Females	255	240	244	244	255	262
60.6 mg/l	Males	259	232	242	254	268	303
	Females	220	203	209	214	222	226
67.9 mg/l	Males	248	231	224	242	264	311
	Females	223	200	208	216	228	241

Necropsy Findings

<u>Group</u>	<u>Finding</u>	<u>Number</u>
Controls	Mottled lungs	8/10
37.9 mg/l	Mottled lungs	2/10
60.6 mg/l	Lung, red foci	2/10
	Mottled lungs	1/10
67.9	Mottled lungs	6/10
	Liver purple	2/10
88.4	Mottled lungs	10/10
	Liver discolored	10/10
	Spleen dark red	1/10
201.9 mg/l	Mottled lungs	10/10
	Lungs blood filled	3/10
	Liver discolored	6/10
Other	Necropsy findings also included gastrointestinal gas distention in animals dying prior to sacrifice. Bladders distended with fluid were also observed in these early deaths.	

Conclusions

Remarks Field

An LC_{50} of 68.4 mg/l was determined using a series of concentrations. Solvent induced narcosis appeared as a common effect and may have been the cause of death at the high concentration. Lung and liver abnormalities were common and severity was dose related.

Data Quality

• Reliability

Klimisch Code 2 Study design, conduct and reporting are considered reliable to address the test endpoint although not conducted in accord with glp standards.

References

An acute inhalation toxicity study of C-12 1 [1,3-Dioxolane] in the rat. Project No. 79-7304, Bio/dynamics Inc., July 11, 1980.

Other

This study is supported by a literature study that reported an LC_{50} of 118 mg/l in the rabbit (exposure time not given), 87 mg/l/ (4 hours) in the male rat and 166 mg/l (4 hours) in the guinea pig.'

References for supporting data

1. Czajkowska, T, Krysiak, B and Popińska, E. Experimental studies of toxic effects of 1,3,5-trioxane and 1,3-dioxolane. I. Acute toxic effect. Med Pr; Vol 38, 1987, P184-90.

Repeated Dose Toxicity, Two-week Gavage

Type Repeated Dose Toxicity, Two-week Inhalation

Test Substance 1,3 -Dioxolane
CAS Number: 646-06-0
Purity listed as 99.9%

Method

- **Guideline**
 - Toxic Substances Control Act Test Guidelines - 798.4900
 - Meets OECD 4 12 Guideline except for lack of clinical chemistry determinations
- **GLP** Yes
- **☑** 1991
- **Species** Rat
- **Strain** CrI:CD[®](SD)BR, Charles River
- **Route of administration** Gavage in corn oil.
- **Duration of Test** Fifteen days
- **Doses** Water control, vehicle control, 75, 250, 750 and 2000 mg/kg/day
- **☑** Rats of each sex.
- **Frequency of Treatment** Daily for fourteen consecutive days.
- **Number of Animals/group** Ten of each sex
- **Control Group and Treatment group** Water treated controls and corn oil treated controls. (Ten animals of each sex per group)
- **Post-Exposure Observation Period** Overnight

- Statistical Methods
 - 0 All hematology data were statistically compared to both control groups using a one-way Analysis of Variance Test and then Dunnett's Test, to determine the statistical significance of the individual groups in comparison with each of the control groups
 - 0 All body weight, feed consumption and organ weight data were statistically compared against both control groups using the following statistical tests: Analysis of Variance, Dunnett's Test, Bartlett's Test of homogeneity, **Kruskal-Wallis** Test and Dunn's Test.
-
- Remarks Field for Test Conditions
- 0 Age at study initiation
 - Males: approximately nine weeks
 - Females: approximately nine weeks
 - 0 Number animals of each sex per dose
 - Ten of each sex per concentration
 - 0 Satellite groups
 - None
 - 0 Housing
 - Individual wire-bottom cages
 - 0 Clinical observations performed and frequency
 - Mortality twice daily and gross signs daily about 30 minutes after dosing. Body weights daily. Feed consumption weekly.
 - 0 Terminal observations
 - Blood samples were evaluated for the following parameters: red blood cell count, hematocrit, hemoglobin, mean corpuscular volume, white blood cell count, white blood cell differential count, red blood indices for mean corpuscular hemoglobin and mean corpuscular hemoglobin concentration, reticulocyte counts (absolute), red blood cell morphological assessment and platelet counts (absolute).
 - Postmortem examination included gross lesions of the thoracic, abdominal and pelvic viscera. organ weights were recorded for the liver, kidneys, adrenals, spleen, pancreas, lungs, pituitary, thymus, as well as the reproductive organs (male rats ▪ testis, epididymides and accessory sex glands; female rats ▪ ovaries and uterus)

- 0 Histopathology
 - Histopathology was performed on all tissues showing gross changes, liver, kidneys, adrenals, spleen, pancreas, lungs, pituitary, thymus, as well as the reproductive organs (male rats - testis, epididymides and accessory sex glands; female rats - ovaries and uterus). Initially, tissues from both controls and the high-dose were examined. Because this preliminary examination of tissues from both control groups and the 2000 **mg/kg/day** dosage group revealed potential histopathology in the kidney, liver, thymus (male and female rats) and testes (male rats), these tissues were also examined in the remaining dosage groups.
- ◊ Differences from OECD Guideline
 - This study is in accord with the current OECD 4 12 guideline for 14 or 28-day inhalation studies with the exception that clinical chemistry parameters were not determined. The current study uses twice the number of animals required by the OECD guideline and one additional dose level.

Results

- NOAEL
 - 0 Males: 75 **mg/kg/day**
 - 0 Females: 75 **mg/kg/day**
- LOEAL
 - 0 Males: 250 **mg/kg/day**
 - 0 Females: 250 **mg/kg/day**
- Mortality
 - 0 Males: Three high dosage group male rats were found dead on days **4, 9** and 14 of the study. Deaths were preceded by clinical observations, and two of these rats had severe weight losses. Necropsy of one rat revealed gastric erosions. No other deaths occurred during the conduct of the study.
 - 0 Females: Four 2000 **mg/kg/day** dosage group female rats died. These deaths occurred on days **3, 8, 9** and 10 of the study and were preceded by clinical observations. These rats had moderate, although sometimes transient, weight losses occur, and necropsy of one rat revealed reddened areas in the stomach. Two additional high dosage group deaths occurred on days 10 and 13 of the study; these deaths were related to intubation accidents. No other deaths occurred during the conduct of the study.

- Toxic Responses

Dose (mg/kg/day)	Toxic Response (statistical comparisons noted are against the corn oil control)
75	<ul style="list-style-type: none"> ■ Males: No significant effects . Females: No significant effects
250	<ul style="list-style-type: none"> . Males <ul style="list-style-type: none"> o Slight reduction in platelets ($p>.05$) o Reduction in body weight gain o Reduced feed consumption . Females <ul style="list-style-type: none"> o Reduction in lymphocyte count ($p<.01$)
750	<ul style="list-style-type: none"> ■ Males <ul style="list-style-type: none"> o Reduction in platelets ($p\leq.01$) o Reduction in body weight gain o Reduced feed consumption o Liver and lung relative weights increased o Reduced relative thymus weight ■ Females <ul style="list-style-type: none"> o Excess salivation in some females. o Reduction in lymphocyte ($p\leq.05$) and reticulocyte count ($p\leq.01$) o Liver relative weight increased o Spleen and thymus relative weights decreased

2000	<ul style="list-style-type: none"> ■ Males <ul style="list-style-type: none"> o Clinical observations after dosing included hypotonia, excess salivation, ataxia, decreased motor activity, rales, impaired and lost righting reflex, labored breathing, red penile exudate and urine-stained abdominal fur o Reduced feed consumption o Reduction in platelets ($p \leq 0.1$) o Overall loss in body weight. o Liver, lung and kidney relative weights increased o Prostate, spleen and thymus weights decreased o Histopathological findings in liver (6/10), kidney (4/10), thymus (6/10) and testis(2/10). ■ Females <ul style="list-style-type: none"> o Clinical observations after dosing included excess salivation (750 and 2000 mg/kg/day), hypotonia, ataxia, decreased motor activity, impaired and lost righting reflex, gasping, bradypnea, oral exudate, rales, lacrimation and urine-stained abdominal fur o Reduction in lymphocyte count ($p \leq 0.1$) o Reduction in platelet count ($p \leq 0.1$) o Reduction in body weight gain o Liver weight marginally increased o Kidney relative weight increased o Spleen weight marginally decreased o Thymus weight decreased o Pancreas relative weight decreased o Histopathological findings in liver (5/10), kidney (3/10), thymus (5/9) and spleen (2/10)
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Remarks Field for 0 Body
Results Weights
Gains

Dose (mg/kg)	Mean Body Weight Gains (grams)			
	Males		Females	
	Day 1-8	Day 8-15	Day 1-8	Day 8-15
Water only	29.5	33.8	11.0	12.6
Corn Oil	28.5	38.5	8.6	11.4
75	27.5	37.4	11.5	9.5
250	26.7	32.6	5.2	10.8
750	20.5	31.0	4.2	16.0
2000	-29.0	9.3	-10.1	11.0

- 0 Clinical Signs
 - o Males: Clinical observations after dosing included hypotonia, excess salivation, ataxia, decreased motor activity, rales, impaired and lost righting reflex, labored breathing, red penile exudate and urine-stained abdominal fur.
 - o Females: Clinical observations attributable to the 750 and 2000 **mg/kg/day** dosages of the test substance included excess salivation (750 and 2000 **mg/kg/day**), hypotonia, ataxia, decreased motor activity, impaired and lost righting reflex, gasping, bradypnea, oral exudate, rales, lacrimation and urine-stained abdominal fur (2000 **mg/kg/day**).

- 0 Hematology
 - o Group designations are used to describe the hematology results, groups are: Group I, water control; Group II, corn oil control; Group III, 75 **mg/kg/day**; Group IV, 250 **mg/kg/day**; Group V, 750 **mg/kg/day**; Group VI, 2000 **mg/kg/day**.
 - o Among male animals, statistically significant differences were seen only in the platelet counts. Groups IV, V, and VI differed significantly from group I at the 0.01 level. Groups V and VI differed significantly from group II at the 0.01 level, but the difference between groups II and IV was not statistically significant

- 0 Hematology
- Among female animals, statistically significant differences were seen in the total leukocyte count, lymphocyte count, platelet count, nucleated red blood cells, and reticulocyte count. In the total leukocyte count, groups IV, V, and VI differed significantly from group I at the 0.05 level; none of the treated groups differed significantly from group II. With respect to lymphocytes, groups IV, V, and VI differed significantly from group I at 0.01; group IV and VI differed significantly from group II at 0.01 while group V differed significantly from group II at 0.05. No significant differences occurred among groups with respect to neutrophils or minor cell types indicating that the differences in total leukocyte count was a function of differences in lymphocytes. No corresponding effect was evident in males. Nucleated red blood cells were significantly different in groups III, V, and VI from group I at 0.05 but did not differ significantly from group II. Reticulocytes were significantly higher in group VI than in group I at 0.05, and lower in group IV than in group II at 0.01. No other differences among groups were significant with respect to reticulocytes. It appears that the differences in nucleated red blood cells and reticulocytes, cells of the same lineage differing only with respect to maturity, are a function of random variation rather than a treatment effect, particularly because differences from controls occur in both directions. In female animals platelets in group VI were significantly lower than in group I and group II at the 0.01 level.
 - From the above data, it appears that a decrease in circulating platelets was a treatment effect as it occurred at a statistically significant levels in both males and females. In males, rats in groups IV, V, and VI had significantly lower platelet counts than group I and groups V and VI had significantly lower platelet counts than group II. In females group VI animals had significantly lower platelet counts than groups I and II. Among the other groups there was a non-significant dose-related trend.
- 0 Hematology Comments
- Treatment related hematologic changes observed in this study were a decrease in platelets, seen in both sexes, and a decrease in lymphocytes (also reflected in total leukocytes) seen in females only. No hematologic treatment effects were observed in the rats given 75 mg/kg/day of the test compound.

V Organ Weights	Treatment related organ weight changes are given above under toxic effects. Organ weigh changes were not remarkable although some reached statistical significance.
V Necropsy findings	Examination of each animal during necropsy did not indicate any exposure-related gross pathologic changes
0 Histopathology	<p>No treatment-related microscopic changes were observed in any of the male and female rats given 75,250, or 750 mg/kg/day of 1,3-Dioxolane.</p> <p>Treatment-related microscopic changes were observed in the liver, kidney, thymus, and testis of rats given 2000 mg/kg/day of 1,3Dioxolane. The treatment -related changes consisted of centrilobular. hepatocellular hypertrophy and midzonal hepatocellular vacuolation (male and female rats); thymic atrophy (male and female rats); renal cortical tubular basophilia and dilatation and accumulations of birefringent intratubular crystals (male rata); subacute renal pyelitis (male and female rats), and multifocal testicular degeneration. Incidence values are given above.</p>

Conclusions

Remarks Field

Gavage administration of Dioxolane to rats of each sex was associated with hematologic effects, body weight gain reductions and organ weight changes at doses of 750 and 2000 **mg/kg/day**. The 250 **mg/kg/day** dose level was associated with decreases in platelets in male rats and lymphocyte counts in female rats.

The authors of the laboratory report elected to call 250 **mg/kg/day** a NOEL in the female rat. The writer of this summary is not in agreement **with** that conclusion and has selected 75 **mg/kg/day** as the appropriate NOEL.

Data Quality

- Reliability

Klimisch Code 1. May be used without restriction.

References

Repeated Dose Oral Toxicity Study of **1,3-Dioxolane** Administered Via Gavage to **CrI:CD[®](SD)BR** Rats. Argus Research Laboratories, Inc. 905 Sheehy Drive Horsham, Pennsylvania 19044, Laboratory Project ID **508-002P**.

Other

- 0 This study is supported by a two-week inhalation study of Dioxolane conducted by **BioDynamics** for Celanese Corporation. In this study Charles River CD rats (**5/sex/group**) were exposed for six hours per day, **five** days per week, for two weeks at concentrations of 0,984 or 3280 ppm. All animals survived the duration of the study. No treatment related observations were recorded. Body weights, organ weights, clinical chemistry parameters, and macropathology were unremarkable in treated animals. Depressed leukocyte values in male and female exposed animals were reported.'
- This study is supported by a **12-day** inhalation study of Dioxolane conducted by Dow Chemical. The primary effect of Dioxolane inhalation exposure to rats was a reduction in white blood cells counts at 23 19 or 5 132 ppm. Body weight gain was affected at the **5132-ppm** exposure level. There was no corresponding myeloid toxicity or inflammatory **foci**².

References for
supporting studies

1. A two week inhalation toxicity study of C- 12 1 (**1,3-Dioxolane**) in the rat. **Bio/Dynamics Inc** Project No. 80-7429, June 22, 1981.
2. **1,3-Dioxolane: A two-Week Vapor Inhalation Toxicology Study** in Fischer 344 Rats. Dow Chemical Company 1989 Study ID K-010634-005

Repeated Dose Toxicity, Four-week Drinking Water

Type Repeated Dose Toxicity, Four-week Oral

Test Substance **1,3-Dioxolane**
CAS Number: 646-06-o

Method

- Guideline None
- GLP No
- Year 1977
- Species Rat, mouse, golden Syrian hamster.
- Strain Charles River strain
- Route of administration Drinking water
- Duration of Test Four weeks
- Doses 0, 0.5, 1.0 and 2.0% in drinking water
- Sex Male and Female
- Exposure Period Twenty-four hours per day
- Frequency of Treatment Continuous, seven days a week.
- Number of Animals/group Five of each sex
- Control Group and Treatment Five animals of each sex exposed only to water
- Post-Exposure Observation Period None
- Statistical Methods Body weight data analyzed by ANOVA followed by Tukey's or Scheffe's test of multiple comparison.

Remarks Field for Test Conditions

◇ Age at study initiation	Not specified
◇ Number animals of each sex per dose	Five of each sex per concentration
◇ Satellite groups	None

- 0 Housing Individual
- 0 Clinical observations performed and frequency
 - . Mortality daily.
 - Body weights weekly.
 - Water consumption twice weekly.
- 0 Terminal observations Body weights
- 0 Test Substance Prepared weekly in tap water.

Results

- NOAEL


- 0 Rats
 - . Males: 0.5%
 - Females: 0.5%
- 0 Mice
 - Males: 0.5%
 - . Females: 0.5%
- 0 Hamsters
 - . Males: 1.0 %
 - . Females: 1.0%

- LOEAL

- 0 Rats
 - . Males: 1.0%
 - Females 1.0%
- 0 Mice
 - Males: 1.0%
 - . Females 1.0%
- 0 Hamsters
 - . Males: 2.0%
 - . Females 2.0%

- Mortality All animals survived the duration of the study.

- Toxic Responses

Dose	Toxic Responses		
0.5%	Reduced water consumption for female rats only.		
1.0%	▪ 	Males	Reduced water consumption Reduction in body weight gain ^{†*}
		Females	Reduced water consumption Slight reduction in body weight gain

	■ Mice.	Males	Slight reduction in body weight gain
		Females	Reduced water consumption Slight reduction in body weight gain
	■ Hamsters	Males	None
		Females	None
2.0%			
	■ Rats	Males	Reduced water consumption Reduction in body weight gain ^{††}
		Females	Reduced water consumption Reduction in body weight gain [†]
	■ Mice	Males	Slight reduction in body weight gain
		Females	Reduced water consumption Slight reduction in body weight gain
	■ Hamsters	Males	Slight reduction in body weight gain
		Females	Slight reduction in body weight gain
	† = p≤ 0.05 †† = p≤ 0.01		
	* Although absolute body weights were not significantly reduced, body weight gain was.		

Remarks Field for
Results

◇ Body
Weights

Mean Body Weights(n=5)					
Drinking Water Level (percent)	Rats				
	Body Weight on Week (grams)				
Males	0	1	2	3	4
0	210	271	331	376	391
0.5	209	264	314	351	364
1.0	210	256	298	332	334
2.0	209	237	273 [†]	304 [†]	309 ^{††}
Females					
0	156	182	209	231	238
0.5	157	178	203	221	227
1.0	156	182	196	208	215
2.0	157	164	182	198 [†]	193 [†]
† = p ≤ 0.05 †† = p ≤ 0.01					

◇ Body Weights

Drinking Water Level (percent)	Mice Body Weight on Week (grams)				
Males	0	1	2	3	4
0	27	33	34	37	38
0.5	28	32	34	36	37
1.0	28	32	35	36	37
2.0	28	30	32	33	34
Females					
0	23	27	28	29	30
0.5	23	26	28	28	28
1.0	23	26	27	27	29
2.0	23	24	25	26	27

◇ Body Weights

Drinking Water Level (percent)	Hamsters Body Weight on Week (grams)				
Males	0	1	2	3	4
0	69	80	93	99	104
I 0.5	69	84	96	104	107
1.0	70	77	90	99	105
2.0	69	77	85	92	95
Females					
0	64	71	80	88	93
0.5	63	74	86	94	100
1.0	64	74	83	90	98
2.0	64	72	79	83	86

● Clinical Signs

None recorded

Conclusions

Remarks Field

- . Rats of each sex showed reduced body weight gains at the two highest dose levels. Body weight gains of high-dose mice and hamsters also appear to be affected but did not achieve statistical significance. The authors of the report only noted the statistically significant changes in body weight gains for rats (males 1.0 and 2.0 %, females 2.0%). Hamsters appear to be less sensitive to the toxic effects of dioxolane than rats and mice and rats appear to be the most sensitive based solely on body weight gains.
- The dosing for the male rats at 1% drinking water with water consumption at 30 ml per day calculates to approximately 1000 **mg/kg/day** Dioxolane. This is the approximate dose level where body weight changes are observed in other studies.

Data Quality

• Reliability

Klimisch Code 2. Reliable with restrictions. Study design, conduct and reporting are considered reliable to address the test endpoint although not conducted in accord with GLP standards.

References

Four-week pilot study with Dioxolane in drinking water of albino rats, albino mice and golden Syrian hamsters. Industrial BIO-Test Laboratories, Inc., Project No. 8560-10579, July 12, 1977.

Other

This study is supported by a 14-day corn-oil gavage study in rats of each sexing which hematologic effects, body weight gain reductions and organ weight changes were reported at doses of 750 and 2000 **mg/kg/day**. The **250-mg/kg/day** dose did not affect body weight gain.

References for supporting studies

Repeated Dose Oral Toxicity Study of **1,3-Dioxolane** Administered Via Gavage to **CrI:CD®(SD)BR** Rats. Argus Research Laboratories, Inc. 905 Sheehy Drive Horsham, Pennsylvania 19044, Laboratory Project ID **508-002P**.

Repeated Dose Toxicity, Two-week Inhalation

Type	Repeated Dose Toxicity, Two-week Inhalation
Test Substance	1,3-Dioxolane CAS Number: 646-06-o Purity 99.99 % by analysis
Method	
• Guideline	This study was conducted to meet the requirements of the United States Toxic Substances Control Act (TSCA), Good Laboratory Practice Standards in 40 CFR part 792; TSCA Health Effects Test Guideline in 40 CFR 798.250; and OECD ISBN 92-64-12367-9 Paris 1982. Study also meets OECD 412 Guideline.
• GLP	Yes
• Year	1989
• Species	Rat
• Strain	Fischer 344 Charles River, Kingston NY
• Route of administration	Whole body inhalation of vapor
• Duration of Test	Twelve days
• Doses	0, 5, 16, 23, 19 or 5, 132 (Mean, measured concentrations)
• Sex	Male and Female
• Exposure Period	Six hours per day
• Frequency of Treatment	◇ Five days a week ◇ Nine treatments
• Number of Animals/group	Five of each sex
• Control Group and Treatment	Five animals of each sex exposed only to air under the same chamber conditions
• Post-Exposure Observation Period	Overnight

- Statistical Methods
 - Descriptive statistics (mean and standard deviation), were used to report chamber concentrations, temperature, and relative humidity.
 - Remaining parameters were first tested for equality of variance using Bartlett's test. If the results for Bartlett's test rejected the equality of variances, the parameter was flagged for careful evaluation of results. All parameters were then subjected to appropriate parametric analysis as described below. In-life body weights, hematologic (excluding differential WBC) and clinical chemistry parameters, terminal body weights, and organ weights (absolute and relative, except testis) were evaluated using a two-way analysis of variance (**ANOVA**) with the factors of sex and dose. Results for absolute and relative testis weights were analyzed using a one-way **ANOVA**. If significant dose effects were determined in the one-way **ANOVA**, then separate doses were compared to controls using Dunnett's test. For those parameters examined by a two-way **ANOVA**, examination was made first for a significant sex-dose interaction. If this existed, a one-way **ANOVA** was done separately for each sex. If no sex-dose interaction was identified, and a dose effect was identified, or if in the subsequent **ANOVAs** separated by sex a dose effect was identified, the separate **ANOVAs** were used for each exposure group with control. To control for multiple comparisons with control, a Bonferroni correction was used.

Remarks Field for
Test Conditions

- ◇ Age at study initiation
 - . Males: eight weeks
 - . Females: eight weeks
- Number animals of each sex per dose
 - . Five of each sex per concentration
- ◇ Satellite groups
 - . None
- ◇ Housing
 - . Not specified in report
- ◇ Clinical observations performed and frequency
 - . Mortality and gross signs daily prior to exposure.
 - . Body weights on days 1, 3, 5, 8, and 11.
 - . Behavior pattern and nervous system activity were assessed by specific observations for lethargy, tremors, convulsions, salivation, lacrimation, diarrhea, and other sign of altered nervous system function.

- ① Terminal observations
 - Blood taken for hematology (red cell count, hemoglobin, hematocrit, total white blood cell count, platelets, differential leukocyte counts and morphology of **WBCs**, **RBCs** and platelets) and clinical chemistry (urea nitrogen, **alanine** aminotransferase activity, aspartate aminotransferase activity, alkaline phosphatase activity, glucose, total protein, albumin, globulin, total bilirubin, cholesterol, triglycerides, phosphorus and calcium)
 - Complete gross postmortem examination.
 - Organ weights for brain, heart, lungs, liver, kidneys, and testis were recorded for all animals.

- ① Histopathology
 - The following tissues were examined from all control and high-exposure animals only. Adrenals, aorta, bone, bone marrow, brain (cerebrum, brainstem, cerebellum), cecum, cervix, coagulating glands, colon, duodenum, epididymides, esophagus, eyes, gross lesions, heart, ileum, jejunum, kidneys, **lacrimal/hardarian** glands, larynx, liver, lungs, mammary gland, mediastinal lymph node, mediastinal tissues, mesenteric lymph node, mesenteric tissues, nasal tissues, oral tissues, ovaries, oviducts, pancreas, parathyroid glands, peripheral nerve, pituitary, prostate, rectum, salivary glands, seminal vesicles, skeletal muscle, skin, spinal cord (cervical, thoracic, lumbar), spleen, stomach, testes, thymus, thyroid-gland, tongue, trachea, urinary bladder, uterus, vagina
 - Microscopic examination of tissues **from** rats in lower **1,3-dioxolane** exposure groups was not conducted since no target organs were identified in the high-exposure group.

- ◇ Differences from OECD Guideline

This study is in accord with **the** current OECD 412 guideline for 14 or 28-day inhalation studies. There are slight differences in the suggested parameters for clinical chemistry and in the duration of the study (**12-days**) but these are not considered scientifically significant.

Results

- NOAEL
 - Males: 516 ppm
 - Females: 516 ppm
- LOEAL
 - Males: 2319 ppm
 - Females: 2319 ppm

- Mortality
 - All animals survived the duration of the study.

- Toxic Responses

Dose	Response
516 ppm	<ul style="list-style-type: none"> ■ No significant effects
2319 ppm	<ul style="list-style-type: none"> • Reduction in white-blood cell counts for males and females. • Decrease in spleen weight in females ■ Increased relative liver weight in females
5132 ppm	<ul style="list-style-type: none"> ■ Reduction in white-blood cell counts for males and females. • Decreased body weight gains for males (11%) and for females (4%) ■ Decreased alertness and slight incoordination at the end of each exposure lasting 45 to 85 minutes. • Other changes detailed below were not considered toxicologically significant.

Remarks Field for Results

◇ Body Weights

Body Weight on Day (grams)					
Males	1	3	5	8	11
Control	136	146	154	166	184
516 ppm	135	146	150	168	176
2319 ppm	134	142	150	165	175
5132 ppm	135	134	139	156	164
Females					
Control	112	117	118	124	129
516 ppm	113	118	120	129	132
2319 ppm	114	118	119	128	131
5132 ppm	112	110	111	121	124

- Clinical Signs

Animals exposed to 5132 ppm dioxolane appeared to have decreased alertness and incoordination at the end of each exposure lasting 45 to 85 minutes.

- Hematology WBC

Mean White Blood Cells ($\times 10^3$)		
Conc (ppm)	Males	Females
Control	9.5 \pm 0.9	7.0 \pm 2.1
516 ppm	9.1 \pm 0.8	7.0 \pm 1.7
2319 ppm	6.9 \pm 0.9	5.2 \pm 0.7
5132 ppm	5.8 \pm 1.4	4.6 \pm 1.2

0 Hematology Comments	<p> . The mean RBC, HGB, HCT and PLAT values of males and females exposed to 23 19 and 5 132 ppm were statistically increased when compared to mean control values. These slight increases were exposure-related but were not considered toxicologically significant. Historical control hematologic data from previous 2-week inhalation studies conducted in this laboratory with Fischer 344 rats revealed that each of the identified differences were within the historical normal range except PLAT values for the 23 19 and 5 132 ppm, females and the 5132 ppm males. </p> <p> . Mean WBC counts of males and females from the two highest exposure groups were statistically decreased from control values in an apparent concentration-dependent manner. Examination of mean WBC differential data indicated increased neutrophil:lymphocyte ratios of male and female rats in these two exposure groups. The WBC counts in this study were within the range of historical controls for 29 two-week inhalation studies in this laboratory. </p> <p> . There was no demonstrable microscopic evidence of inflammatory foci in tissues or decreased cellularity of lymphoid organs and bone marrow. </p>
0 Clinical Chemistry	<p> Clinical chemistry analyses indicated statistically decreased mean values for AP in the 23 19 and 5 132 males, and the 5 132-ppm females when compared to respective control means. These slightly decreased values were not considered toxicologically significant. </p>
0 Organ Weights	<p> The relative weights of brain, heart, kidneys, liver, lungs, and testes of male rats exposed to 5 132 ppm were statistically increased from those of male controls. Similarly, relative weights of heart, liver, and lungs from high exposure group females were increased statistically from the control females. All of the relative organ weight changes were secondary to decreased terminal body weights. </p>
0 Necropsy findings	<p> Examination of each animal during necropsy did not indicate any exposure-related gross pathologic changes </p>
0 Histopath- ology	<p> The few microscopic changes seen during the histopathologic examination were considered to be spontaneous events unrelated to exposure. There were no microscopic changes in the brains of high-exposure group rats to account for the transitory clinical signs observed after each exposure.. </p>

Conclusions

Remarks Field

The primary effect of dioxolane inhalation exposure to rats was a reduction in white blood cells counts at 23 19 or 5 132 ppm. Body weight gain was affected at the 5 132-ppm exposure level. There was no corresponding myeloid toxicity or inflammatory foci.

Data Quality

• Reliability

Klimisch Code 1. May be used without restriction.

References

1,3-Dioxolane: A two-Week Vapor Inhalation Toxicology Study in Fischer 344 Rats. Dow Chemical Company 1989 Study ID K-O 10634-005

Other

This study is supported by a two-week inhalation study of Dioxolane conducted by **BioDynamics** for Celanese Corporation. In this study Charles River CD rats (**5/sex/group**) were exposed for six hours per day, five days per week, for two weeks at concentrations of 0,984 or 3280 ppm. All animals survived the duration of the study. No treatment related observations were recorded. Body weights, organ weights, clinical chemistry parameters, and macropathology were unremarkable in treated animals. Depressed leukocyte values in male and female exposed animals were reported.'

References for supporting studies

A two-week inhalation toxicity study of C-121 (**1,3-Dioxolane**) in the rat. **Bio/Dynamics** Inc Project No. 80-7429, June 22, 1981.

Repeated Dose Toxicity, **13-Week Inhalation**

Type Repeated Dose Toxicity, **13-Week Inhalation**

- Test Substance **1,3-Dioxolane**
CAS Number: 646-06-0
Purity >**99.78** % by analysis

Method

- Guideline This study was conducted to meet the requirements of the United States Toxic Substances Control Act (TSCA), Good Laboratory Practice Standards in 40 CFR part 792; TSCA Health Effects Test Guideline in 40 CFR 798.250; and OECD ISBN 92-64-12367-9 Paris 1982.
- GLP Yes
- Year 1990
- Species Rat
- Strain Fischer 344
Charles River, Kingston NY
- Route of administration Whole body inhalation of vapor
- Duration of Test Thirteen weeks plus eight week recovery study
- Doses 0,298, 1000 or 3010 (Mean, measured concentrations)
- Sex Male and Female
- Exposure Period Six hours per day
- Frequency of Treatment Five days a week
Thirteen weeks
- Number of Animals/group Ten of each sex
- Control Group and Treatment Ten animals of each sex exposed only to air under the same chamber conditions
- Post-Exposure Observation Period 24 Hours for main groups; eight weeks for satellite recovery groups.

- Statistical Methods
 - Descriptive statistics (mean and standard deviation), were used to report chamber concentrations, temperature, and relative humidity.
 - Body weights, absolute and relative organ weights, clinical chemistry, bone marrow **myeloid/erythroid** ratios and appropriate hematology data for animals to be terminated after 13 weeks were evaluated by Bartlett's test for equality of variances. Clinical chemistry, bone, marrow **myeloid/erythroid** ratios and appropriate hematology data for satellite animals were evaluated the same way.
 - WBC differential counts are normally analyzed, in this laboratory, with descriptive statistics. Because effects on the white cell was anticipated based on a two-week study, absolute lymphocyte numbers, absolute neutrophil numbers and myeloid to erythroid cell ratios were evaluated in the following statistical evaluation. Based on the outcome of Bartlett's test, exploratory data analysis was performed by a parametric or non-parametric analysis of variance (ANOVA), followed respectively by **Dunnett's** test or the **Wilcoxon** Rank-Sum test with a Bonferroni correction for multiple comparisons.
 - Statistical outliers were to be identified by a sequential test and excluded from analysis only for documented scientifically sound reasons that were unrelated to exposure.

Remarks Field for Test Conditions

- Age at study initiation
 - Males: seven weeks
 - Females: seven weeks
- Number animals of each sex per dose
 - Ten of each sex per concentration
- Satellite groups
 - Ten of each sex per dose level and control were exposed for thirteen weeks. Blood samples were taken at four and thirteen weeks of exposure and again at four and eight weeks into the recovery period. Satellite rats were necropsied at the end of the eight-week recovery period.
- Housing
 - Not specified in report
- Clinical observations performed and frequency
 - Mortality and gross signs daily
 - Body weights: weekly
 - Functional observation battery during the sixth and last week of exposure for main group animals consisting of observations for any unusual conditions with respect to: pupil size, respiration, movement, skin and haircoat, salivation, lacrimation, urine staining, fecal staining, locomotor behavior and responsiveness to touch, noise and tail pinch.
 - Animals were evaluated for alertness and activity at the beginning and end of each exposure to evaluate sedative effects

Results

- NOAEL **0** Males: 1000 ppm
0 Females: 298 ppm
- LOEAL **0** Males: 3010 ppm
0 Females: 1000 ppm
- Mortality All animals survived the duration of the study.

Toxic Responses

Dose	Response
298 ppm	. No significant effects
1000 ppm	. Reduction in white-blood cell counts for females. . Decrease in spleen weight in females . Increased relative liver weight in females
3010 ppm	. Reduction in white-blood cell counts for males and females. ▪ Mild changes in liver of males. ▪ Decreased spleen weights in males and females ▪ Decreased alertness at the end of each exposure . Decreased urine specific gravity

Remarks Field for Results

◇ Body Weights

	Exposure Period Day (n=20)			Recovery Period Day (n=10)		
Males	Day-1	46	86	96	116	152
Control	150	253	299	294	325	323
298 ppm	150	253	300	294	319	314
1000 ppm	147	247	298	287	315	311
3010 ppm	147	257	306	303	332	329
Females						
Control	109	164	182	186	192	187
298 ppm	108	165	183	180	191	186
1000 ppm	108	161	182	183	196	190
3010 ppm	106	158	179	173	189	188

- **Clinical Signs** ● Some exposed male and female rats had notable clinical observations including: red swollen eyes, perineal soiling, diarrhea and darkened crusty material around the eyes and/or nares. One male rat exposed to 1000 ppm had persistent redness, swelling and/or crusty material around one eye. This condition, however, was resolved by study termination. The orbital sinus blood sampling procedure may have resulted in some of the observations of crusty material around the eyes. Because of a lack of a dose response relationship and persistence in any of the observations, none were considered to be exposure-related or toxicologically significant
- Animals exposed to **3000-ppm** dioxolane appeared to have decreased alertness and responsiveness at the end of each exposure. The animals appeared to be fully recovered by the time they were removed from the exposure chamber (approximately 30 minutes after cessation of test material exposure),
- **Functional Observation Battery** Functional Observational Battery exams were conducted at 6 and 13 weeks. Isolated observations of perineal soiling, decreased activity (one observation) and chromorhinorrhea were noted. These were not thought to be exposure-related due to the low incidence and lack of a dose response.

● **Hematology**

◇ **WBC**

Conc (ppm)	Mean White Blood Cells ($\times 10^3$)			
	Week of Study			
Males	4-week	13-week	Recov-4	Recov-8
Control	10.7 \pm 2.1	9.1 \pm 1.3	10.2 \pm 1.0	10.0 \pm 1.2
298	9.4 \pm 2.1	9.6 \pm 1.4	8.2 \pm 0.8*	9.1 \pm 1.1
1000	8.2 \pm 0.8*	8.6 \pm 1.0	8.7 \pm 1.5*	8.4 \pm 0.5*
3010	5.5 \pm 1.0*	6.0 \pm 1.0*	8.9 \pm 0.9*	8.6 \pm 1.2*
Females				
Control	7.3 \pm 1.4	9.2 \pm 0.9	7.2 \pm 1.3	6.4 \pm 1.1
298	8.7 \pm 1.6*	9.1 \pm 1.1	6.4 \pm 1.1	6.7 \pm 0.8
1000	6.2 \pm 0.8	7.8 \pm 0.5*	6.1 \pm 0.9	6.0 \pm 0.6
3010	4.9 \pm 0.9*	5.1 \pm 0.6*	6.0 \pm 1.0*	5.8 \pm 0.6

- **Hematology**

Comments

 - Although exposure decreased the WBC of rats the magnitude of the decrease at any given time interval was never below the lower limits of historical controls.
 - The percent eosinophils in 3010 ppm exposed rats was increased relative to controls after 13 weeks of exposure
 - Small increases in mean platelet counts occurred during the exposure period in some of the 1000 and 3010 ppm groups. There was a single observation of increased platelets in females exposed to 300 ppm at 4 weeks, however, there was no apparent exposure-response relationship and no difference at **13-weeks**. The minimal statistical platelet elevations in the 1000 and **3000-ppm** groups were within available historical control data; hence these differences were considered to be of minor biological significance.
 - Sporadic statistically significant variances in red blood cells and hematocrit were in the range of historical controls and did not show concentration-response relationships and are considered of no toxicological significance

- **Clinical Chemistry**

A number of statistically identified differences in various clinical chemistry parameters in exposed male or female rats were identified when compared to their respective controls; however, there was no clear relationship to exposure. Since these parameters were generally within historical control ranges or were not associated with corresponding gross or microscopic effects, most of these clinical chemistry changes were not considered to be diagnostic of a target organ effect. Increased ALT in 3000 ppm exposed rats at 4 weeks, and, in females only at 13 weeks, could be related to the small increases in liver weight. Histological changes were also noted in livers of 3010 ppm exposed males.

- ◇ **Urinalysis**

Male and female rats exposed to 3010 ppm for 13 weeks (main study and satellite groups) had statistically decreased urine specific gravity. Repeat urinalyses revealed similar results. The specific gravity alteration was not associated with a morphologic indication of nephrotoxicity. It may represent a physiologic response to repeated exposure to dioxolane vapors.

I

Organ Weights

0

0

0

O

Necropsy findings

Cr

- Histopathology
 - Except for liver changes in high-dose male rats, there were no pathologic findings of significance. In male rats exposed to 3000 ppm dioxo lane, hepatocytes in centrilobular regions of lobules were slightly larger and had more cytoplasmic eosinophilia than controls (incidence 1 0/ 10).
 - Microscopic examination of bone from exposed and control rats following 13 weeks of exposure indicated that myeloid and erythroid cells were approximately similar in number and, had normal morphology and apparently normal maturation sequences. In addition, there was no cytotoxicity of the microenvironment at the light microscopic level. Therefore, the decreases in WBC counts were not associated with observable histopathologic changes in bone marrow.
 - The ratio was lower in male and female rats exposed to 3010 ppm for 13 weeks relative to controls; however, the **myeloid/erythroid** ratio of this exposure group was similar to that of respective controls after 8 weeks post exposure.

Conclusions

Remarks Field The primary effect of dioxolane inhalation exposure to rats was a reduction in white blood cells counts at 1000 or 3000 ppm. Rats appeared healthy, body weight gain was not affected and there was no corresponding myeloid toxicity although there was a slight reduction in myeloid cells of the bone marrow at 3000 ppm. Exposure related pathologic effects were limited to slight enlargement of centrilobular hepatocytes of males at 3000 ppm. There was no indication of testicular toxicity based on careful light microscopic examination.

Data Quality

- Reliability Klimisch Code 1. May be used without restriction.

References

1,3-Dioxolane: 13-Week Vapor Inhalation Toxicology Study in Fischer 344 Rats. Dow Chemical Company 1990

Other

This study is supported by a **12-month** inhalation study in the literature where a concentrations of 2500 **mg/m³** was found not to produce dominant lethal effects or significant signs of toxicity manifest by gross observations or mortality'

References for supporting studies

¹ **Barański** B; Stetkiewicz J; Czajkowska T; Sitarek **K**; Szymczak W Mutagenic and gonadotoxic properties of trioxane and dioxolane Med Pr; Vo135: P245-55 (1984)

* = p < 0.05

Reverse Mutation Assay - *S. typhimurium*

Type Reverse mutation assay - *S. typhimurium*

Test Substance **1,3-Dioxolane**
CAS Number: 646-06-0

Method

- **Guideline** None specified
- **GLP** No
- **System of Testing** Bacterial
- **Year** 1980
- **Species/Strain** *S. typhimurium*: TA98, TA100, TA1535, TA 1537, TA1538
- **Metabolic activation**
 - Tested with and without
 - Rat liver S-9
 - Aroclor 1245 induced
 - 0.5 ml S-9 per 100 ml agar plate
- **Concentrations tested**
 - TA98 0, 0.005, 0.010, 0.10, 1, 5, 10, 25, 50 µl/plate
 - ◇ TA100 0, 0.005, 0.010, 0.10, 1, 5, 10, 25, 50 µl/plate
 - ◇ TA1535 0, 0.005, 0.010, 0.10, 1, 5, 10 µl/plate
 - ◇ TA1537 0, 0.005, 0.010, 0.10, 1, 5, 10 µl/plate
 - ◇ TA1538 0, 0.005, 0.010, 0.10, 1, 5, 10 µl/plate
 - All tested with and without metabolic activation
- **Statistical Methods**

Evaluation criteria were used as follows:

 - Strains TA-1535, TA-1537 and TA-1538: If the solvent control value is within the normal range, a test material that produces a positive dose response over three concentrations with the highest increase equal to three times the solvent control value will be considered to be mutagenic.
 - ◇ Strains TA-98 and TA-100: If the solvent control value is within the normal range, a test material that produces a positive dose response over three concentrations with the highest increase equal to twice the solvent control value for TA-98 and TA-100 will be considered to be mutagenic.

- Remarks Field for Test Conditions
- ◊ One replicate
 - ◊ Positive Controls
 - o Without activation
 - Sodium azide: TA100, TA 1535
 - . 9-Aminoacridine: TA1537
 - . 2-Nitrofluorene: TA98, TA153 8
 - o With activation
 - . 2-Anthramine: all strains
 - V Solvent: none
 - V Repeat study: TA-1535 was repeated with and without activation because of low solvent values observed in the initial assay.
 - V Evaluation Criteria: Dose-response related increase in revertants specific for historical rates in each strain

Results

- Result No mutagenic activity under activation or non-activation conditions.
- Cytotoxic Concentration No significant cytotoxicity at any concentration
- Genotoxic Effects
 - . Not genotoxic under non-activation conditions
 - . Not genotoxic under activation conditions.

- Remarks Field for Results
- o Material was soluble in water

Conclusions

- Remarks Field
- o Not genotoxic under non-activation conditions
 - o Not genotoxic under activation conditions.

Data Quality

- Reliability Klimisch Code 2. Reliable with restrictions. Study design, conduct and reporting are considered reliable to address the test endpoint. Study was conducted under glp's. Conditions of the assay concerning time of incubation may vary somewhat from current OECD guideline.

References

Mutagenicity Evaluation of C-1 2 1 (1,3-Dioxolane) in the Ames Salmonella/Microsome Plate Test, Litton Bionetics Study 20988, 2/29/1980 submitted to Celanese Chemical Company.

Other

This study is supported by other bacterial genotoxicity studies specifically:

- 0 A negative single-plate **5-strain** Salmonella reverse mutations assay with and without metabolic activation by Goodyear'
- 0 A negative 5-strain Salmonella reverse mutations assay with and without metabolic activation by Hilltop conducted for Ferro under glp conditions.*
- 0 Published negative **5-strain** Salmonella reverse mutations assay with and without metabolic **activation**.³
- 0 A combined *S. typhimurium* and *Saccharomyces cerevisiae* mutation evaluation conducted by Litton Bionetics for Dow Chemical **Company**.⁴

References for Supporting Studies

1. Mutagenicity Evaluation of **1,3-Dioxolane**, Goodyear Tire and Rubber Company. Laboratory report **#79-55, 29** November 1979.
2. The **Salmonella/Microsomal** Assay for Bacterial Mutagenic Activity of Unstabilized **1,3-Dioxolane**. Hilltop Research Project No 83-0212-21 10 March, 1983.
3. Kowalski Z, Spiechowicz E, **Barański** B. Absence of mutagenicity of trioxane and dioxolane in *Salmonella typhimurium*. **Mutat Res** 136: 169-71 (1984).
4. Mutagenicity Evaluation of Compound **D8 (1,3 -Dioxolane)**. Litton Bionetics, 1/20/1975 submitted to Dow Chemical Company.

Mutagenic Assay - *S. typhimurium* and *Saccharomyces cerevisiae*

Type	Mutagenic assay - <i>S. typhimurium</i> and <i>Saccharomyces cerevisiae</i>	
Test Substance	1,3-Dioxolane CAS Number: 646-06-0	
Method		
• Guideline	None specified	
• GLP	No	
• System of Testing	Bacterial	
• Year	1975	
• Species/Strain	<i>S. typhimurium</i> TA1535, TA 1537, TA1538 <i>Saccharomyces cerevisiae</i> strain D4	
• Metabolic activation	<input type="checkbox"/> Tested with and without induced rat, mouse and monkey homogenates of liver, lung and testis. Animals were induced with a “mixture of polychlorinated biphenyls”.	
• Concentrations tested	<input type="checkbox"/> <i>Saccharomyces cerevisiae</i> : <ul style="list-style-type: none">o 2.0 and 5.0% in suspension <input type="checkbox"/> Salmonella: <ul style="list-style-type: none">o 0.75 and 1.50% for suspension incubationso 1.5% for plate tests. <input type="checkbox"/> All tested with and without metabolic activation	
• Statistical Methods	None specified	

- Remarks Field for Test Conditions
 - Duplicate plates, one replicate
 - Salmonella were tested with rat tissues in the suspension test only. Mouse and monkey tissues were used in the plate incorporation technique only.
 - Salmonella and yeast were tested at high and low dose levels using **non-**activation conditions and activation conditions with rat liver, rat lung and rat testis homogenates.
 - Salmonella were tested under activation conditions with mouse liver, mouse lung and mouse testis homogenates only at the high dose level.
 - Salmonella were tested under activation conditions with monkey liver, monkey **lung** and monkey testes homogenates only at the high dose level.
 - Yeast were tested only with rat tissue activation and not mouse or monkey.
 - Positive Controls
 - Without activation
 - . Ethylmethane sulfonate
 - . 2-Nitrofluorene
 - . Quinacrine mustard
 - With activation
 - 2-Acetylfluorene
 - . Dimethylnitrosamine
 - Solvent: none (except for positive controls)
 - Evaluation Criteria: Dose-response related increase in revertants specific for historical rates in each strain.

Results

- Result
 - No mutagenic activity under activation or non-activation conditions.
- Cytotoxic Concentration
 - Fifty percent bacteria survival when tested at 3 .0% in the preliminary cytotoxicity test, yeast showed no cytotoxicity at 5% (highest concentration tested).
- Genotoxic Effects
 - . Not genotoxic under non-activation conditions
 - . Not genotoxic under activation conditions.

Remarks Field for Results

- . With the organ homogenates, the positive controls gave significant genotoxic activity only with the liver and testis preparation for the rat, the liver for the mouse and the liver for the monkey.
- . Under non-activation conditions, both bacteria and yeast responded as expected to the positive controls.

Conclusions

Remarks Field

- . Not genotoxic under non-activation conditions
- . Not genotoxic under activation conditions.
- The contribution of the lung and testis homogenates to the genotoxic evaluation of the test substance is questionable since the positive controls did not show significant response except for the rat testis with strain TA-1537
- . The main value of this study is the negative response of the yeast as other more robust bacterial genotoxicity tests have been conducted using this test material.

Data Quality

• Reliability

Klimisch Code 2. Reliable **with** restrictions. Study design, conduct and reporting are considered reliable to address the test endpoint although the study was not conducted under glp's.

References

Mutagenicity Evaluation of Compound D8 (**1,3-Dioxolane**). Litton Bionetics, 1/20/1975 submitted to Dow Chemical Company.

Other

The Salmonella portion of this study is supported by other Salmonella tests providing clear negative results. The key study is listed below.

References for Supporting Studies

Mutagenicity Evaluation of C- 12 1 (**1,3-Dioxolane**) in the Ames **Salmonella/Microsome** Plate Test, Litton Bionetics Study 20988, 2/29/1980 submitted to Celanese Chemical Company

***In Vitro* Mammalian Chromosome Aberration Test**

Type *In Vitro* Mammalian Chromosome Aberration Test

Test Substance **1,3-Dioxolane**
CAS Number: 646-06-0

Method

- **Guideline** **None** specified
- **GLP** **Yes**
- **System of Testing** **Non-bacterial**
- **Year** **1985**
- **Species/Strain** **Chinese Hamster Ovary (CHO-WBL), Origin S. Wolff's laboratory UCSF, Litton Bionetics clone, average cell cycle time 12-14 hours.**
- **Metabolic activation** **Tested with and without**
 - o **Rat liver S-9**
 - o **Aroclor 1245 induced**
 - o **0 15 microliters per 10 ml culture**
- **Concentrations tested** **0, 2.0, 3.0, 4.0, 5.0 mg/ml for both non-activation and activation conditions.**
- **Statistical Methods** **Fishers Exact Test for comparisons of treated cultures with pooled controls.**

- Remarks Field for ◇ Test Conditions
- Range-finding test to determine cytotoxicity or cell cycle delay used the following conditions:
- o Single culture per dose levels of 0, 1.67 or 5.0 **mg/ml** for both **non-activation** and activation conditions. Level of 0.50 was added under activation conditions.
 - o Under non-activation conditions there was a 2-hour exposure to test substance followed by 23 hours of incubation prior to addition of colcemid and 2.5 hours of incubation prior to collection.
 - o Under activation conditions the same time intervals were employed and fetal-calf serum was omitted.
- The aberration test used the following conditions:
- o Duplicate cultures at each dose level of **0, 2.0, 3.0, 4.0, 5.0 mg/ml** for both non-activation and activation conditions
 - o Solvent, McCoy's 5a culture media.
 - o Under non-activation conditions, there was a **7.5-hour** exposure followed by 2.5-hr incubation in the presence of colcemid.
 - o Under non-activation conditions there was a 2-hour exposure to test material followed by a wash and an **8-hour** incubation in the presence of colcemid.
 - o 100 Metaphases from each culture were scored for a total of 200 metaphases per dose level.
 - o Negative control, McCoy's 5a
- 0 Positive controls
- . Without activation – Mitomycin C
 - . With activation – Cyclophosphamide

Results

- Result

No increase in the number of aberrations was found at any concentration of test material. Positive controls demonstrated the sensitivity of the test system.
- Cytotoxic Concentration

Cytotoxicity was not observed up to and including the highest concentration tested of 5 **mg/ml** which is the highest concentration recommended under the current OECD 473 guideline.
- Genotoxic Effects
 - Not genotoxic under non-activation conditions
 - Not genotoxic under activation conditions.

Remarks Field for Results

- Material was soluble in water
- . The number of aberrations per cell, the percent cells with aberrations, and the percent cells with more than one aberration were all similar to appropriate controls under both activation and non-activation conditions.
- . Positive controls produced clear significant increases in aberrations as measured by aberrations per cell, the percent cells with aberrations and the percent cells with more than one aberration.

Conclusions

Remarks Field

- Not genotoxic under non-activation conditions
- . Not genotoxic under activation conditions.
- Study was well conducted, although the incubation times and exact procedures vary somewhat from the current OECD guideline the consistently low level of aberrations observed at all dose levels both with and without activation and the clear responses of the positive controls strongly support lack of genotoxic activity in this assay.

Data Quality

- Reliability

Klimisch Code 2. Reliable with restrictions. Study design, conduct and reporting are considered reliable to address the test endpoint. Study was conducted under glp's. Conditions of the assay concerning time of incubation vary somewhat from current OECD guideline.

References

Mutagenicity Evaluation of **1,3-Dioxolane** in an In Vitro Cytogenic Assay Measuring Chromosome Aberration Frequencies In Chinese Hamster Ovary (CHO) Cells. Submitted to Ferro Corporation. Litton Bionetics, Inc, Kensington Maryland. LBI Project No. 20990, February 1985

Mouse Lymphoma Forward Mutation Assay

Type Mouse lymphoma forward mutation assay

Test Substance 1,3-Dioxolane
CAS Number: 646-06-0

Method

- **Guideline** None specified
- **GLP** No
- **System of Testing** Non-bacterial
- **Year** 1985
- **Species/Strain** Mouse lymphoma L5 178Y TK+/-
- **Metabolic activation** Tested with and without
 - Rat liver S-9
 - Aroclor 1245 inducedFinal concentration of S-9 in cell suspensions was five percent.
- **Concentrations tested** 0,750, 1500, 2500, 3000, 4000, 50000 nl/ml for both non-activation and activation conditions.
- **Statistical Methods** Simple ratio criteria

Remarks Field for Test Conditions

- ◇ Initial trial under activation conditions discarded because the positive control mutation frequencies were unacceptable.
- ◇ Duplicate plates for counting mutant colonies
- ◇ Solvent, water
- ◇ Negative control, water
- ◇ Positive controls
 - Without activation – EMS (2 levels)
 - With activation – MCA (2 levels)

Results

- **Result** No dose-dependent increase in the number of mutants in the absence or presence of metabolic activation was observed. Dioxolane did not induce significant increases in the mutation frequency at the TK locus in L5178Y TK⁺/– cells. . Positive controls demonstrated the sensitivity of the test system.
- **Cytotoxic Concentration** Low dose-dependent cytotoxicity observed under non-activation conditions percent relative growths ranges from 85 % to 61% . Under activation conditions, cytotoxicity was dose-dependent but low, with percent relative growths ranging from 95% to 78%.
- **Genotoxic Effects**
 - Not genotoxic under non-activation conditions
 - Not genotoxic under activation conditions.

Remarks Field for Results

- Material was soluble in water
- Under non-activation conditions, the mutant frequency ranged from 12.8 to 22.5 x 10⁻⁶ without any discernable dose dependency.
- Under activation conditions, the mutant frequency ranged from 26.5 to 43.9.5 x 10⁻⁶ without any discernable dose dependency
- The minimum mutation frequency required for a positive response under non-activation conditions based on the control mutation frequencies in this assay was 38 X 10⁻⁶.
- The minimum mutation frequency required for a positive response under activation conditions, based on the control mutation frequencies in this assay, was 46 X 10⁻⁶.
- The discarded initial activation assay had positive control values that were considered unacceptably low, although clearly positive. In this discarded test, the mutation frequency for the test substance was also low and similar to that found in the repeated test. The discarded test supports the final result.

Conclusions

Remarks Field

- 0 Not genotoxic under non-activation conditions
- 0 Not genotoxic under activation conditions.
- 0 Study was well conducted, although there was no glp certification the study appears to have been conducted using a glp-quality protocol in a glp compliant laboratory.

Data Quality

- Reliability

Klimisch Code 2. Reliable with restrictions. Study design, conduct and reporting are considered reliable to address the test endpoint although not conducted in accord with glp standards.

References

Mutagenicity Evaluation of 1,3-Dioxolane in the Mouse Lymphoma Forward Mutation Assay. Submitted to Ferro Corporation. Litton Bionetics, Inc, Kensington Maryland. LBI Project No. 20989, April 1985

Other

Mammalian Cell Transformation Assay, *in vitro*

Type	Mammalian Cell Transformation Assay, <i>in vitro</i>
Test Substance	1,3-Dioxolane CAS Number: 646-06-0
Method	
• Guideline	None specified
• GLP	No
• Year	1981
• Cell Type	C3H 10T-1/2 clone 8
• Concentrations tested	0, 1, 10, 20, 50, 100, 500, 1000, 2000, 5000, 10000, 20000 and 40000 µg/ml
• Statistical Methods	None used

Remarks Field for Test Conditions	<p>◇ Dose selection based on initial cytotoxicity testing using plating efficiency as a metric.</p> <ul style="list-style-type: none"> ● A set of replicates was exposed in T-25 flasks and a set of replicates was exposed in flat 60 mm culture dishes. ● Incubation of plates carried out in sealed Mason jars to prevent volatilization ● T-25 flasks were sealed to prevent loss of test material. ● Twenty-four hour incubation time with test substance ● Cells washed after 24-hour incubation with test substance ● Cells fed with fresh media 24 hours after exposure was complete and every three days until the 14-day point then weekly (with 10%FCS supplementation) for the duration of the 38 day culture period. ● At the 14-day interval, four to six T-25 flasks at each concentration and three to five dishes were stained and examined. ● At the 38-day interval, two to four T-25 flasks at each concentration and three or four dishes were stained and examined. ● High-concentration levels (>2000 µg/ml) of T-25 flasks at the 38-day interval were contaminated with a fungus in the initial assay and were run a second time about 2 months later. In this repeat assay, six T-25 flasks were incubated as before for 38 days at each concentration level from 1 to 20,000 µg/ml, stained and examined. The data varied from the initial assay in that there was a much higher transformation rate in the positive controls (BP 5 or 6 times higher) and the transformation rate for the test substance was much higher at the repeated concentration levels. The negative control still showed no transformed colonies. ● Positive control: Benzpyrene @ 2.5, 1.0 or 0.5 µg/ml ● Colonies examined macroscopically and microscopically. ● 300 cells plated per flask or dish <p>◇ Differences from Guideline The study is similar to the Draft OECD Guideline “<i>In Vitro</i> Syrian Hamster Embryo Cell Transformation Assay”. The major differences from this guideline are the type of cells used and the incubation times employed. It was similar or superior in other respects including number of dose levels employed.</p>
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Results

- Result
 - Report authors conclude that Dioxolane is a “weak positive” in this cell-transformation assay.
 - The data suggest transforming activity but there are issues of concern with the data. The poor reproducibility, the lack of clear dose-concentration relationships, the test material sometimes giving results similar to positive controls and at other times no response, and the lack of high-concentration toxicity data make it difficult to conclude that this assay is positive.
 - A more conservative interpretation of the data is “Not interpretable”

- **Cytotoxic Concentration** Cytotoxicity observed above 5,000 **ug/ml**. There was no valid measure of cytotoxicity given for the **38-day** examinations

- **Genotoxic Effects** Not interpretable

Remarks Field for Results 0 The majority of transformed foci were classified as “type II” rather than “type III” This is defined as follows:

- Type I. Foci composed of monolayer cells that are more densely packed than the background cells. This type is not considered malignant and is not scored.
- Type II. Foci show massive piling up into virtually opaque multilayers. The cells are only moderately polar, thus criss-crossing is not pronounced. Fifty percent of Type II foci have been shown to be malignantly transformed.
- Type III. Foci are composed of highly polar, fibroblastic, multilayered, criss-crossed arrays of densely stained cells. Eighty-five percent of Type III foci have been shown to be malignantly transformed.

	14-Day Foci/plate		38-Day Foci/plate		Repeat 38-day
Conc (ug/ml)	T-25	Dishes	T-25	Dishes	T-25
0	0.0	0.0	0.0	0.0	0
1	0.0	0.0	0.0	0.0	2.8
10	0.0	0.0	0.0	0.0	2.8
20	0.0	0.0	0.0	0.0	3.7
50	0.0	0.0	0.0	0.0	3.7
100	0.0	0.0	1.0	0.75	4.7
500	0.0	0.0	0.3	1.33	4.0
1,000	nn	0.0	1.5	1.25	5.0
2,000	0.6	.	C		
5,000	0.0	0.0 [†]	C	2.0	4.66
10,000	0.5 [†]	0.0 [†]	C	2.25	1.2
20,000	0.0 ^{††}	0.0 [”]	C	0.25	0
40,000	-	0.0 ^{††}	.	.	
BP 0.5	0.5	0.75	0.5	6	3.0
BP 1.0	1.0	1.0	1.25	12	6.0
BP 2.5					7.0
- = Not run, [†] = Cytotoxic, ^{††} = Highly Cytotoxic C = contaminated, BP = Benzpyene Positive Control					

Conclusions

Remarks Field

The report authors concluded that these results represent a weak positive; however, the reviewer (who prepared this summary) disagrees with this interpretation and judges the study not interpretable with regard to genotoxic activity.

Data Quality

- Reliability

Klimisch Code 3. Reliable with severe restrictions, study design, conduct and reporting are considered reliable to address the endpoint but the data are not clearly supportive of the author's conclusions.

References

An Assay of Cell Transformation and Cytotoxicity in **C3H10T1/2** Clonal Cell Line for the Test Chemical C-121 (**1,3-Dioxolane**). University of Minnesota Environmental Pathology Laboratory, Authors Garry and Nelson, 27 August 1981.

Other

This study is supported by a study using **BALB/C-3T3** cell cultures. This study, however, reported Dioxolane to give a clear negative response in cell transformation.

Reference for supporting study

Evaluation of 1,3-Dioxolane in the In Vitro Transformation of **BALB/C-3T3** Cells Assay. Litton Bionetics, Inc, Project 20992, January 1985. Submitted to Ferro Corporation

Mammalian Cell Transformation Assay, *in vitro*

Type	Mammalian Cell Transformation Assay, <i>in vitro</i>
Test Substance	1,3-Dioxolane CAS Number: 646-06-o
Method	
• Guideline	None specified
• GLP	Not specified (report was reviewed by the quality assurance unit but did not specifically state that the test was conducted under glp)
• 中国标准	1985
• Cell Type	◇ BALB/C-3T3 ◇ Subclone 14 of Clone I-13 from Takeo Kakunaga
• Concentrations tested	0, 1000, 4000, 8000, 12000 and 15000 nl/ml
• Statistical Methods	Bailey's modification of Student's t-test was applied to calculate the statistical significance of the positive control and test material transforming activity.
Remarks Field for Test Conditions	◇ Positive control: 2.0 micrograms/ml MCA ◇ Seventy-two hour treatment time followed by washing and continued incubation for four weeks with re-feeding twice a week. ◇ Treatments carried out in sealed 25 cm ² flasks to prevent volatilization. ◇ Doses based on preliminary cytotoxicity range-finding test. ◇ 30,000 cells plated per culture vessel ◇ 18 Cultures per dose level (lower n in Results table due to contaminated flasks) ◇ Simultaneous cytotoxicity test conducted under same conditions as transformation assay to verify the toxicity of test material to cells.

Preliminary Cytotoxicity Concentration (nl/ml)	Average Number of Colonies per Flask	Relative cell Survival (%)
0	94.3	100.0
1000	92.0	97.6
2000	86.3	91.6
4000	74.0	78.5
8000	55.0	58.3
16000	4.0	4.2
Simultaneous I Cytotoxicity Test		
0	59.7	100.0
1000	51.0	85.4
4000	46.3	77.6
8000	29.7	49.7
12000	8.3	14.0
15000	0.0	0.0

- ◇ Differences from Guideline The study is similar to the draft OECD Guideline “*In Vitro* Syrian Hamster Embryo Cell Transformation Assay”. The major differences from this guideline are the type of cells used and the incubation times employed. It was similar or superior in other respects including number of dose levels, replicates, cytotoxicity testing and MTD selection. The current study protocol was shown to be sensitive by the response of the positive controls.

Results

- Result No increase in the number of transformed colonies or transformed foci was noted at any concentration of test material. Positive controls demonstrated the sensitivity of the test system
- Cytotoxic Concentration Severe cytotoxicity observed above 12000 micrograms/ml, moderate dose-dependent cytotoxicity observed at 1000 microgram per ml and above. Data in table above
- Genotoxic Effects Not genotoxic under these conditions

Remarks Field for Results The test material was water soluble and precipitation was not recorded.

Results	Transforminn Activity		
Concentration (nl/ml)	Log ₁₀ (x+1) (mean±sem)	(n)	Foci/Culture average
0	0.050±.027	(18)	0.12
1000	0.050±.027	(18)	0.12
4000	0.067±.030	(18)	0.17
8000	0.050±.027	(18)	0.12
12000	0.067±.030	(18)	0.17
15000	0.075±.034	(16)	0.19
MCA (2ug/ml)	0.602±.039	(18)	3.00**
** p<0.01; MCA = 3 Methylcholanthrene			

Conclusions

Remarks Field

No genotoxic activity. The transforming activities of all five doses of Dioxolane were comparable to the spontaneous background average rate of 0.12 foci/culture vessel and there was no evidence of a dose-related response with the test material. This range of test material treatments corresponded to 0 to 85% survival in the simultaneous colony survival assay. Although the survival assay, using only 200 cells, suggested there would be no surviving cells with test material treatments at 15000 nl/ml, some cells survived in the transformation assay where 30,000 cells were plated per culture vessel. Therefore, concentrations of 1,3-dioxolane from 1,000 to 15,000 nl/ml were evaluated as being nontransforming to Balb/C-3T3 cells.

Data Quality

- Reliability

Klimisch Code 2. Reliable with restrictions, study design, conduct and reporting are considered reliable to address the test endpoint although not conducted in accord with glp standards.

References

Evaluation of 1,3-Dioxolane in the In Vitro Transformation of BALB/C-3T3 Cells Assay. Litton Bionetics, Inc, Project 20992, January 1985. Submitted to Ferro Corporation

Other

This study is supported by a previous study using C3H 10T-1/2 cell cultures. This study reported Dioxolane to give a weak positive response in cell transformation but is considered uninteruptible by the writer of the robust summary.

Reference for Supporting Study

An Assay of Cell Transformation and Cytotoxicity in C3H 1 OT 1/2 Clonal Cell Line for the Test Chemical C-121 (1,3-Dioxolane). University of Minnesota Environmental Pathology Laboratory, Authors Garry and Nelson, 27 August 1981

Mouse Micronucleus Assay

Type Mouse micronucleus assay

Test Substance **1,3-Dioxolane**
CAS Number: 646-06-o

Method

- Guideline None specified, Meets or exceeds OECD 474 guideline
- GLP Yes
- Year 1989
- Species/Strain Mice/ICR Harlan Sprague Dawley
- Sex Mice of each sex
- Route of Administration IP Injection in Corn Oil
- Doses 0, 525, 1050 and 2100 mg/kg
- Exposure Period 24, 48 and 72 hours
- Statistical Methods The incidence of micronucleated polychromatic cells was compared to controls with a significant increase relative to the negative control set at the $p \leq 0.05$ level using Kastenbaum-Bowman Tables*

Remarks Field for Test Conditions

◇ Age at Study Initiation Six to eight weeks

◇ Number of animals per dose Five of each sex at each dose-time sacrifice were planned, more animals were dosed due to mortality. High-dose mortality was unexpectedly high and a supplemental group of 15 animals of each sex was dosed at 2100 mg/kg. Of these supplemental animals only four of each sex survived to the scheduled sacrifice at 72 hours. There were insufficient surviving high-dose animals to sacrifice five per dose at the high dose. The resulting dosing and sacrifice of animals is shown below:

	Number of Animals Sacrificed at Each Time					
Dose	24 hours		48 hours		72 hours	
(mg/kg)	M	F	M	F	M	F
0	5	5	5	5	5	5
525	5	5	5	5	5	5
1050	5	5	5	5	5	5
2100	5	5	3	5	4	4

-
- 0 Control Groups and Treatment
 - Negative Control: Corn Oil
 - Positive Control: Triethylenemelamine in water at 0.25 mg/kg
 - 0 Clinical Observations
 - All low and mid-dose animals appeared normal. High-dose animals showed several clinical observations indicating toxicity including prostration, lethargy, irregular breathing, and ruffled fur.
 - 0 Criteria for Evaluating Results
 - Negative control must not exceed an incidence of 5/1000 micronucleated polychromatic cells.
 - A positive result is indicated if the incidence of micronucleated polychromatic cells is significantly increased relative to the negative control ($p \leq 0.05$, Kastenbaum-Bowman Tables)
 - 1000 polychromatic erythrocytes scored for each animal.
 - 0 Criteria For Selection of MTD
 - The high dose was set as 80% of the LD₅₀ (7 day). A preliminary toxicity study was conducted with results shown below. The LD₅₀ (7 day) was calculated by probit analysis to be approximately 2603 mg/kg.

Prelim Tox Study	Seven Day Mortality	
IP Dose (mg/kg)	Males	Females
Contol	0/5	0/5
1576	0/5	0/5
2048	1/5	0/5
2663 I	5/5	3/5 I
3462	5/5	5/5
4500 I	4/5	5/5 I

- 0 Differences from Current OECD Guideline

This study meets all requirements of the current OECD 474 guideline. The study used more dose levels and time points than indicated in the guideline. As there was a previous positive study in the literature the study design was especially robust.

Results

- Effect on PCE/NCE ratio by dose level by sex.

	PCE/Total Erythrocytes					
Dose	24 Hours		48 Hours		72 Hours	
(mg/kg)	M	F	M	F	M	F
0	0.59	0.59	0.62	0.62	0.62	0.66
525	0.53	0.57	0.62	0.64	0.54	0.65
1050	0.53	0.60	0.58	0.67	0.43	0.54
2100	0.31	0.44	0.26	0.41	0.37	0.35
TEM	0.57	0.57	-	-	-	-

- Genotoxic Effects Negative

- NOAEL 1050 mg/kg for bone marrow toxicity as measured by the PCE/NCE ratio

Remarks Field for Results

◇ Induction of Micronucleated Cells

	Micronucleated Cells/ 1000 PCE					
Dose	24 Hours		48 Hours		72 Hours	
(mg/kg)	M	F	M	F	M	F
0	0.2	0.2	0.2	0	0.2	0
525	0.2	0.2	0.2	0	0.4	0.6
1050	0.2	0.2	0	0.4	0	0
2100	0.6	0	0.3	0.4	0	0
TEM	18.4	20.8	-	-	-	-

- Mortality
 - All low and mid-dose mice survived to sacrifice.
 - Mortality at high dose was males 23/35 and females 21/35

Conclusions

Remarks Field

- 0 There was no induction of micronucleated polychromatic cells associated with administration of Dioxolane.
- 0 There was significant induction of micronucleated polychromatic cells associated with administration of the positive control.
- 0 Dioxolane showed no genotoxicity in this assay under these conditions.
- 0 The highest dose of dioxolane was toxic to the animals including the bone marrow.

Data Quality

- Reliability

Klimisch Code 1. Reliable without restriction.

References

Micronucleus Cytogenetic Assay in Mice. Test article C-121 [1,3-Dioxolane]. Microbiological Associates Inc., Study No. T9034.122, December 28, 1989

* Kastenbaum, M and K Bowman. Tables for determining the statistical significance of mutation frequencies. *Mutation Res.* **9:527-549** (1970)

Other

An earlier study in the literature reported Dioxolane was positive in the micronucleus assay. In this report mice were given dioxolane ip in two doses at 24 hr intervals. Dioxolane at doses from 1500 **mg/kg** to 6000 **mg/kg** caused a significant increase of micronucleated polychromatic erythrocytes in bone marrow of mice as compared to the negative control value.

Reference for Supporting Study

Przybojewska B, Dziubautowska E, Kowalski Z. Genotoxic effects of dioxolane and trioxane in mice evaluated by the micronucleus test. *Toxicol Lett* **21:349-52** (1984)

Dominant Lethal, Oral Dosing

Type Dominant Lethal, Oral Dosing

Test Substance Dioxolane
CAS Number: 646-06-0

Method

- Guideline None specified but basically in accord with OECD 478
- GLP No
- Year 1984
- Species Rat
- Strain Albino, Wistar
- Route of administration Oral gavage
- Doses 0, 580, 1160 mg/kg
- Sex Male
- Number of Animals/group 10
- Vehicle Water

Remarks Field for Test Conditions

- | | | |
|---|---|---|
| 0 | Age at Study Initiation | 3 ½ to 4 months; weight 300 to 320 g |
| 0 | Doses | 0,580 and 1160 mg/kg/day |
| 0 | Dosing | Males only |
| 0 | Dosing Schedule | Five days a week |
| 0 | Dosing Duration | Eight weeks |
| 0 | Mating Interval | Weekly |
| 0 | Mating Ratio | 2: 1 females:males |
| 0 | Variations from OECD 478 Protocol Guideline | Guideline suggests that the number of males should be sufficient that 30-50 pregnant rats be evaluated at each time interval. In this study, 13 to 20 pregnant females were evaluated at each time; however, more time periods than typical were evaluated in this study. Current guideline suggests three dose levels; only two were used in this study. |

Results

- Result No evidence of dominant lethal effect

- Dominant Lethal Rate

	Week (Dominant Lethals per Female)							
Dose	1	2	3	4	5	6	7	8
0 mg/kg	0.84	0.83	1.45	1.29	0.89	1.11	0.93	0.67
580 mg/kg	0.73	1.31	0.94	0.89	0.75	0.78	1.06	1.29
1160 mg/kg	0.94	0.64	0.88	1.23	0.56	0.46	0.47	0.67

- Number of deaths at each dose level

Dose	Mortality
0 mg/kg	0/10
580 mg/kg	1/10
1160 mg/kg	1/10

Remarks Field for Results

- ◇ Time of death 580 mg/kg group one animal during fifth week
1160 mg/kg group one animal during third week

- ◇ Clinical Signs Behavior and appearance did not differ to any significant degree from controls

- ◇ Body Weights

Dose (mg/kg)	Body Weight gain (percent control gain)
580	47 %
1160	22 %

- ◇ Organ Weights Absolute and relative weights of liver, kidney and spermatoc vesicle were increased at both dose levels.

- ◇ Necropsy and Microscopic Findings Necropsy finding are not discussed. Histopathology was performed on the testes. Necrosis of the seminiferous epithelium was reported in 1/10 controls, number unspecified in the 580 mg/kg group and 3/9 high dose males.

- 0 Target Organs Testis considered target by histopathology but no change observed in fertility. Other organ weights were altered but there is no histopathology for confirmation. No changes in testes weight resulted from dosing.
- 0 Criteria for Dominant Lethal Effect Investigators used early resorptions as a measure of dominant lethal effect; however, neither implants per female, live fetuses per female nor preimplantation loss per female was affected by treatment.

Conclusions

Remarks Field

Study documentation is good for a published article. The study appears to have been well conducted and is similar to current OECD guideline. Based on reduction in body weight gains, organ weight changes and histopathology all dose levels produced clear signs of toxicity indicating the study is valid regarding use of a minimally toxic dose as the high-dose level. There was no evidence of a dominant lethal effect.

Data Quality

• Reliability

Klimisch Code 2. Study design, conduct and reporting are considered reliable to address the test endpoint although not conducted in accord with glp standards. Study design and reporting meets current EPA/OECD guidelines with minor exceptions.

References

Baranski, B; Stetkiewicz, J. Evaluation of Mutagenic and Gonadotoxic Properties of Trioxane and Dioxolane. Medycyna **Pracy**:35 245-255 (1984)

Other

Dominant Lethal, Inhalation

Type	Dominant Lethal, Inhalation		
Test Substance	Dioxolane CAS Number: 646-06-o		
Method	<ul style="list-style-type: none"> Guideline: None specified conducted basically in accord with OECD 478 GLP: No Year: 1984 Species: Rat Strain: Albino, Wistar Route of administration: Whole body inhalation Doses: 0, 2500 mg/m³ Sex: Male Number of Animals/group: 14 Vehicle: Air Duration of Dosing: 12 Months Statistical Methods: Kruskal Wallis Test followed by non-parametric tests for groups with and without normal distribution. 		
Remarks Field for Test Conditions	0 Age at Study Initiation	3 ½ to 4 months; weight 300 to 320 g	
	0 Doses	0 and 2500 mg/m ³	
	0 Dosing	Males only	
	0 Dosing Schedule	Five days a week, five hours a day.	
	0 Dosing Duration	12 Months	
	0 Mating Interval	Once at end of study for one week	
	0 Mating Ratio	2: 1 females:males	

- ◊ Variations from OECD 478 Protocol Guideline Guideline suggests that the number of males should be sufficient that 30-50 pregnant rats be evaluated at each time interval. In this study, 18 to 20 pregnant females were evaluated. Current guideline suggests three dose levels; only one was used in this study.
- 0 Other Concentration of test material measured by chromatography using a published procedure.

Results

- Result No evidence of dominant lethal effect

- Dominant Lethal Rate

Dose	Effect (per female)					
	Number Pregnant	Live Fetuses	Dead Implants	Number Implants	Corpora Lutea	Preimplantation loss
0 mg/m ³	20	11.4	0.6	12.0	13.7	1.8
2500 mg/m ³	18	11.1	0.17	11.3	13.6	2.3

- Number of deaths at each dose level

Dose	Mortality
0 mg/m ³	0/14
2500 mg/m ³	0/14

Remarks Field for Results

- ◊ Time of death No deaths
- 0 Clinical Signs Not reported
- ◊ Body Weights Not reported
- 0 Organ Weights Not reported

◇ Necropsy and Microscopic Findings	Not clearly reported. It was stated that for the “rats exposed to Dioxolane the frequency of necrosis of the spermidal epithelium was much higher than for the control group rats”; but the incidence or other details was not given. Lydig cell lesions were also mentioned but it was also stated that the treated and control groups were not different regarding this lesion.
◇ Target Organs	None reported.
● Criteria for Dominant Lethal Effect	Investigators used early resorptions as a measure of dominant lethal effect; however, neither implants per female, live fetuses per female nor preimplantation loss per female was affected by treatment.

Conclusions

Remarks Field	Study documentation is good for a published article. The study appears to have been well conducted and the protocol is similar to the current OECD guideline. Based on reduction in body weigh gains, organ weight changes and histopathology, all dose levels produced clear signs of toxicity indicating the study is valid regarding use of a maximum tolerated dose. There was no evidence of a dominant lethal effect.
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Data Quality

● Reliability	Klimisch Code 2. Study design, conduct and reporting are considered reliable to address the test endpoint although not conducted in accord with glp standards. Study design and reporting meets current EPA/OECD guidelines with minor exceptions.
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References

Baranski, B; Stetkiewicz, J. Evaluation of Mutagenic and Gonadotoxic Properties of Trioxane and Dioxolane. *Medycyna Pracy*:35 245-255 (1984)

Other

One Generation Reproduction Study (Drinking Water, low dose)

Type One Generation reproduction Study

Test Substance 1,3-Dioxolane
CAS Number: 646-06-o

Method

- Guideline None, generally follows OECD 415
- GLP No
- Year 1976
- Species Rat
- Strain Charles River strain albino rats (Charles River Wilmington Mass)
- Route of administration Oral, drinking water
- Doses 0.01, 0.03 or 0.1 % in drinking water
- Sex Males and females
- Control Group and Treatment Received water without test substance
- Frequency of Treatment Seven days a week
- Duration of Test 21 Weeks
- Premating Exposure Period for Males 90 Days
- Premating Exposure Period for Females None for F 1 a generation
- Statistical Methods Not described other than that a 95% confidence interval was used to evaluate statistical significance.
- Number of Animals/group 10
- Vehicle Water

Remarks Field for Test Conditions	0	Age at Study Initiation	Not specified
	0	Dosing	Dosed water was available seven days a week and 24 hours a day.
	◊	Test Material Preparation	Test material was prepared weekly and added to the water bottle. Bottles were changed weekly.
	0	Dosing Duration	Dosing started 90 days prior to mating for males. Females were dosed starting with the mating period and throughout gestation and lactation.
	●	Mating Interval	Mating continued for 15 days or until conception was confirmed.
	◊	Examinations	Females were examined daily for evidence of copulation.
	◊	Mating Ratio	Two females to one male
	◊	Culling	Litters of more than 10 pups were reduced to that number by random sacrifice on the fourth day of the lactation period.
	●	Weaning	Litters were weaned 21 days postpartum
	●	Variations from current OECD 415 Protocol Guideline	Significant variations include: <ul style="list-style-type: none"> ◦ Number of pregnant females was about 10 rather than about 20. ◦ Females were treated from the beginning of the mating period rather than from two weeks prior to mating. ◦ Necropsies and microscopic evaluations were not conducted on animals.
	0	Comments	The test material is volatile and significant loss occurs from open systems. Experience indicates that it is stable in solution in closed systems. The weekly preparation and changing of water bottles likely resulted in a small loss of test material concentration as the week progressed. It is unlikely that this affects the results.

Results

- Result
 - 0 No treatment-related effects were noted among the indices used to evaluate reproductive performance.
 - 0 No significantly significant differences were noted between the control and test group with regard to the number of pups delivered stillborn or viable or cannibalized and viable at lactation days 1, 4, 12, or 21.
 - 0 No treatment-related effects were noted among the progeny survival indices.
 - 0 No treatment-related effects in body weight were noted for the F1 a progeny at lactation days 0, 1, 4, 12 or 21.
 - 0 No treatment-related body weight differences were noted for the female parental rats.
- NOAEL 0.1% in drinking water for parental and Fla generations

Remarks Field for Results 0 Mortality One gravid high-dose female died during the Fla gestation period. Necropsy examination revealed no gross abnormalities

Dose	Indices for Parental Parameters Fla Litter (percent)				
	Female Mating	Female Fecundity	Male Fertility	Female Fertility	Parturition
0	76.9	100	100	100	100
0.01	60.0	100	100	90	88.9
0.03	47.1	87.5	87.5	70	85.7
0.10	84.6	90.0	100	100	90.0

0 Parental Body Weights Parental body weight gains were unaffected by treatment with the test article.

Dose	Live Birth Index	Survival Indices*		# Pups Weaned		21-day mean Body Weights (g)	
		4-Day	21-Day	Male	Female	Male	Female
0	100	94.9	96.7	46	42	42	40
0.01	97.6	97.5	82.1	35	39	20	39
0.03	96.1	90.5	80.9	26	44	29	42
0.10	100	81.9	94.7	41	40	30	40

- 0 Fla Exam
 - o All delivered pups appeared outwardly normal. An examination of each pup for external abnormalities was conducted at weaning and all pups were judged free of abnormalities.
 - o The significantly reduced mean body weights of 2 1 -day male pups in the 0.01 dose group is considered spurious due to lack of dose response correlation.
- 0 Clinical Signs
 - None noted

Conclusions

Remarks Field

Adverse effects recorded were limited to a significant but likely spurious difference between low-dose treated and control animals in decreased male pup body weight (low-dose group on lactation day 21). No significant differences were observed between treated and control animals in the following: maternal mortality, mating and fertility indices, male and female fertility, incidence of parturition, gross abnormalities in pups, number of pups delivered stillborn or viable, or cannibalized (lactation days 1, 4, 12 or 2 1), survival indices, female pup body weight, and female parental body weight. It is concluded that 0.1% in drinking water represents a LOAEL for parental and F 1 a generations.

Data Quality

- Reliability

Klimisch Code 2. Study design, conduct and reporting are considered reliable to address the test endpoint although not conducted in accord with glp standards.

References

One-Generation Reproduction Study with 1,3-Dioxolane in Drinking Water in Albino Rats Results of the FO Generation. Industrial Bio-Test Laboratories, Inc. Submitted to PPG Industries March 26 1976.

Other

These results are supported by an inhalation study at 125 ppm in which no significant effects on parental or pup parameters were observed.^{1,2}

References for supporting studies

1. Single Generation Reproduction Study with the Vapors of Dioxolane in Albino Rats Results of the FO and F 1 a Generations, Status Report. IBT No. 663-05562 Industrial Bio-Test Laboratories, Inc. Submitted to PPG Industries March 1, 1976.
2. Single Generation Reproduction Study with the Vapors of Dioxolane in Albino Rats Results of the FO and Flb Generations, Status Report. IBT No. 663-05562 Industrial Bio-Test Laboratories, Inc. Submitted to PPG Industries April 13, 1976.

Definitions

- o Female Mating Index = $100(\text{number copulations}^i / \text{number estrus cycles required})$
 - o Female Fecundity Index = $100(\text{number pregnancies} / \text{number copulations})$
 - o Male Fertility Index = $100(\text{number sires} / \text{number males mated})$
 - o Female Fertility Index = $100(\text{number of pregnancies} / \text{number of females mated})$
 - o Parturition Index = $100(\text{number of parturitions} / \text{number pregnancies})$
 - o Survival Index, 4-day = $100(\text{number pups viable at lactation day 4} / \text{number of viable pups born})$
 - o Survival Index, 21-day = $100(\text{number pups viable at lactation day 21} / \text{number of viable pups at lactation day 4})$
- i) Only one copulation counted per estrus cycle.

One Generation Reproduction Study (Drinking Water, high dose)

Type	One Generation reproduction Study with F 1 a and F 1 b litters
Test Substance	1,3-Dioxolane CAS Number: 646-06-0
Method	
● Guideline	None, generally follows OECD 415
● GLP	No
● 1975	1975
● Species	Rat
● Strain	Charles River strain albino rats (Charles River Wilmington Mass)
● Route of administration	Oral, drinking water
● Doses	0.5 or 1.0 % in drinking water
● Males and females	Males and females
● Control Group and Treatment	Received water without test substance
● Frequency of Treatment	Seven days a week
● Duration of Test	25 Weeks
● Premating Exposure Period for Males	90 Days for males used to produce Fla pups, males used to produce Flb pups were untreated.
● Premating Exposure Period for Females	None for Fla generation, dams for Flb generation were treated for approx 67 days prior to mating. This started with the first mating period and throughout gestation, lactation and a 10-day rest period after weaning of F 1 a litter before mating a second time.
● Statistical Methods	Not described other than that a 95% confidence interval was used to evaluate statistical significance.
● Number of Animals/group	10
● Vehicle	Water

Remarks Field for Test Conditions	0	Age at Study Initiation	Not specified
	0	Dosing	Dosed water was available seven days a week and 24 hours a day.
	0	Test Material Preparation	Test material was prepared weekly and added to the water bottle. Bottles were changed weekly.
	0	Dosing Duration	Dosing started 90 days prior to mating for males. Females were dosed starting with the mating period and throughout gestation, lactation and a 1 O-day rest period after weaning of F 1a litter before mating a second time.
	0	Mating Interval	Mating continued for 15 days or until conception was confirmed.
	0	Examinations	Females were examined daily for evidence of copulation.
	0	Mating Ratio	Two females to one male
	0	Culling	Litters of more than 10 pups were reduced to that number by random sacrifice on the fourth day of the lactation period.
	0	Weaning	Litters were weaned 21 days postpartum
	0	Production of F1b Generation Pups	Due to the severe effects on offspring and possible males in the F 1a generation, treated dams were rested for 10 days after F 1a pups were weaned, and mated with unexposed proven breeders. Females were exposed to test substance during the 10-day interval after weaning F 1a pups. Neither dams nor males were exposed after the F 1b mating interval started.
	0	Sacrifice, Gross Pathology and Organ Weights	Following weaning of the F 1b litter, all surviving female rats were necropsied. Early deaths were also necropsied. Weights of adrenal glands, gonads, pituitary glands and uterus were recorded at necropsy. Fixed tissues (adrenals, ovaries, pituitary glands, uterus and all apparent lesions) were sectioned and stained with H&E for microscopic examination.
	0	Variations from current OECD 415 Protocol Guideline	Significant variations include: <ul style="list-style-type: none"> o Number of pregnant females was about 10 rather than about 20. o Females were treated from the beginning of the mating period rather than from two weeks prior to mating in the F1a mating.

0 Comments

The test material is volatile and significant loss occurs from open systems. Experience indicates that it is stable in solution in closed systems. The weekly preparation and changing of water bottles likely resulted in a small loss of test material concentration as the week progressed. It is unlikely that this affects the results especially in light of the significant effects produced by treatment.

Results

• Result

0 Mating with males pretreated 90 days with dioxolane to produce the F la litter:

- o Treated groups copulated less frequently than controls.
- o Fewer pregnant treated animals delivered than controls.
- o One high-dose male failed to impregnate either female he was paired with and was considered sterile
- o There was a decrease in the number of pups delivered by the high-dose group.
- o Both treated groups showed an increase in the number of stillborn pups.
- o There was reduced survival in progeny from treated animals as compared to controls. Especially the high-dose **24-hour** survival, which was zero percent. Reduced survival indices were noted for the 0.5% group at each time interval examined.
- o Dam's body weight was significantly less than control's for the high-dose group on postpartum days 1 and 4.

0 Mating with proven breeders to produce the Flb litter

- o There was a decrease in the fecundity index and in the parturition index in the test groups.
- o There was a lower female fertility index for the exposed groups.
- o Gross necropsy and histopathology of treated dams was unremarkable, although there was a tendency for changes in the ovaries.
- o Male fertility (unexposed males) was 100%
- o Mating index was unaffected
- o All pups appeared normal and no significant differences were noted with respect to number of pups born, stillborn, cannibalized, born viable or at lactation days 1, 4, 12, and 21. Pup body weight was unaffected.

- NOAEL Not found, effects at all concentrations tested for FO and F1 generations
- LOAEL
 - FO (P) Generation: 0.50%
 - F1 a Generation: 0.50%
 - F1 b Generation: Not applicable since no concurrent exposure

Remarks Field for **◇ Mortality** One high-dose male died during the last week of the **90-day** treatment period, necropsy revealed the cause of death due to respiratory problems not related to test article exposure.

0 Parental Data Fla Litter	Indices for Parental Parameters F1a Litter (percent)				
	Dose (%)	Female Mating	Female Fecundity	Male Fertility	Female Fertility
	0	81.8	100	100	90.0
	0.5	40.9	88.9	100	80.0
	1.0	35.3	100	75	75.0

Number pregnant: control=9; 0.5% = 8, 1.0% = 6 (one high-dose male was found to be sterile)

0 Parental Body Weights Dams Fla	Mean weights of dams on day indicated						
	Dose	G-15	Part	PP-1	PP-4	PP-12	PP-21
	0	375	369	364	366	345	352
	0.5	398	362	344	367	377	371
	1.0	359	331	324 ^a	316 ^a	338	338

G-15 = gestation day 15, Part= day of parturition, PP=postpartum day
a = significant at p< 0.05

0 Progeny Data Fla Litter	Number of pups that were			
	Dose	Delivered	Stillborn	Cannibalized
	0	55	2	0
	0.5	45	6	0
	1.0	14	7	1

◇ Fla data	Survival Indices*		# Pups Weaned		21-day mean Body Weights (g)	
	Dose (%)	Live Birth Index	4-Day	21-Day	Male	Female
	0	96.4	92.5	70.2	15	18
	0.5	86.7	61.5	41.7	8	2
	1.0	42.9	0.0	0.0	0	0

- 0 F1a Exam All pups delivered appeared outwardly normal. An examination of each pup for external abnormalities was conducted at weaning and all pups were judged free of abnormalities.

- 0 Clinical Signs None noted

- 0 Mortality During the Flb Mating and Gestation One high-dose female died post-mating. Necropsy showed non-gravid female with pyometra.

0 F1b Litter Data, Parental Data	Indices for Parental Parameters F1b Litter (percent)					
	Dose (%)	Female Mating	Female Fecundity	Male Fertility	Female Fertility	Parturition
	0	64.3	88.9	100	80	100
	0.5	64.3	66.7	80	60	83.3
	1.0	66.7	50.0	75	50	75

- 0 Dam Body Weights Body weights of dams were unaffected by prior treatment with dioxolane during the mating, parturition and lactation phase of the F 1 b litter production

0 Flb Litter Data, Pup Data		Number of pups that were: [total (mean per dam)]			
	Dose	Delivered	Stillborn	Cannibalized	Viable
	0	78 (9.8)	2(0.3)	0(0.0)	76(9.5)
	0.5	43(8.9)	2(0.3)	0(0.0)	41(8.2)
	1.0	24(8.0)	0(0.0)	0(0.0)	24(8.0)

0 Flb Litter Data, Pup Data			Survival Indices*		# Pups Weaned		21-day mean Body Weights (g)	
	Dose (%)	Live Birth Index	4-Day	21-Day	Male	Female	Male	Female
	0	97.4	92.1	86.9	30	23	45	44
	0.5	95.3	92.7	69.4	10	15	57	51
	1.0	100	91.7	72.7	9	9	59	44

- 0 Flb Pup Exam All pups delivered appeared outwardly normal. An examination of each pup for external abnormalities was conducted at weaning and all pups were judged free of abnormalities.

0 Clinical Signs None noted

0 Gross Pathology No significant differences were noted between control and test group parental females upon gross pathologic examination.

0 Dam Organ Weights

Dose	Organ weights, absolute in grams, relative in g/100g bw					
	Adrenals		Ovaries		Uterus	
	abs	rel	abs	rel	abs	rel
0	0.082	0.0208	0.072	0.0179	0.713	0.1803
0.5	0.076	0.0184	0.075	0.0184	0.797	0.1909
1.0	0.071	0.0172	0.065	0.0156	0.686	0.1658

Adrenals, ovaries and uterus weights showed a tendency to reduced weight with increasing dose. None were statistically different from control but the (n) was relatively small. Other organs showed no tendency to decreased weight with increasing dose of dioxolane.

0 Histopathology of Dams 0 All weighed organs were examined microscopically. No histopathological alterations attributable to the effects of the test material were observed. In treated rats, the hilus of the ovary was unusually prominent and the amount of ovarian parenchyma appeared deficient.

0 Although not considered significant by the examining pathologist, the incidence of ovarian focal "leutinizization of stroma" was 0/10, 1/10 and 3/8 for control, low dose and high dose animals respectively.

Conclusions

Remarks Field

Drinking water exposure of rats at 0.5 or 1 .0% produced clear adverse effects on reproduction. In the fla mating, it could not be determined if effects were attributable primarily to exposure of males, females or both. Survival of **high-**dose pups was especially affected. In the second exposure using untreated proven male breeders, the effects were less severe suggesting that there was a male contribution to the reproductive toxicity. This conclusion is clouded, however, since the females were not exposed to test article during mating, gestation or lactation during the production of this second litter. It is clear that the females were still affected and therefore the female was an affected sex regarding reproductive toxicity of dioxolane.

Data Quality

- Reliability

Klimisch Code 2. Study design, conduct and reporting are considered reliable to address the test endpoint although not conducted in accord with glp standards.

References

One-Generation Reproduction Study with Dioxolane in Albino Rats, Results of the FO Generation. Industrial Bio-Test Laboratories, Inc. Submitted to PPG Industries July 21, 1975.

One-Generation Reproduction Study with Dioxolane in Drinking Water of Albino Rats Results of the FO Generation Females and Proven Breeder Males Industrial Bio-Test Laboratories, Inc. Submitted to PPG Industries February 16, 1976.

Other

The perinatal mortality is supported by a the study of prenatal and postnatal development published by Sitarek et al. in which an increased perinatal death rate was reported after dosing dioxolane at 1150 mg/kg/day, every other day, from day 2-20 of gestation.

References for supporting studies



Sitarek K, Baranski B, **Berlinska** B. The effect of maternal exposure to dioxolane on prenatal and postnatal development in rats. *Pol J Occup Med Environ Health* 5:159-66 (1992).

Definitions

- Female Mating Index = $100(\text{number copulations}^i / \text{number estrus cycles required})$
- Female Fecundity Index = $100(\text{number pregnancies} / \text{number copulations})$
- Male Fertility Index = $100(\text{number sires} / \text{number males mated})$
- Female Fertility Index = $100(\text{number of pregnancies} / \text{number of females mated})$
- Parturition Index = $100(\text{number of parturitions} / \text{number pregnancies})$
- Survival Index, 4-day = $100(\text{number pups viable at lactation day 4} / \text{number of viable pups born})$
- Survival Index, 21-day = $100(\text{number pups viable at lactation day 21} / \text{number of viable pups at lactation day 4})$

i) Only one copulation counted per estrus cycle.

One Generation Reproduction Study, Inhalation

Type	One Generation reproduction Study with F 1 a and F 1 b litters
Test Substance	1,3-Dioxolane CAS Number: 646-06-o
Method	
• Guideline	None, generally follows OECD 4 15
• GLP	No
• 	1975
• Species	Rat
• Strain	Charles River strain albino rats (Charles River Wilmington Mass)
• Route of administration	Inhalation, whole body exposure
• Doses	125 ppm
• 	Males and females
• Control Group and Treatment	Untreated controls, under same chamber housing conditions, same number of animals per group
• Frequency of Treatment	Five days a week (seven hour a day exposure)
• Duration of Test	24 Weeks
• Premating Exposure Period for Males	90 Days for males used to produce Fla pups, 120 days for males (different males) used to produce Flb pups.
• Premating Exposure Period for Females	None for Fla generation, dams for Flb generation were treated for approx 35 days during the F 1 a mating and gestation, then remained unexposed for approximately 35 days prior to mating and the second exposure period.
• Statistical Methods	Not described other than that a 95% confidence interval was used to evaluate statistical significance.
• Number of Animals/group	10
• Vehicle	Air

Remarks Field for Test Conditions	0 Age at Study Initiation	Not specified
	0 Dosing	Inhalation, only water was supplied during exposure. Exposures were conducted in the same chamber being used for an ongoing chronic study of Dioxolane at the laboratory.
	0 Test Material Preparation and Measurement	The inhalation chambers were supplied with vapors of dioxolane generated by bubbling nitrogen gas through test material in a gas-washing bottle. These vapors were diluted with filtered-conditioned air to achieve the desired concentration. Vapor concentration was measured by gas chromatography hourly during the first two weeks of the study then twice daily for the remainder of the study.
	0 Dosing Duration	<ul style="list-style-type: none"> o P males for Fla generation were treated starting 90 days prior to mating. These males were sacrificed after mating. o P males for F lb generation were selected from the population in an ongoing chronic study and had been treated for 120 days prior to mating. o Dams for the F 1 a litter production were exposed starting with the 15-day mating period and through gestation. Exposure was stopped a day or two before parturition. o Dams for the Flb litter production were exposed about 35 days previously during F 1 a litter production. They were then not exposed during delivery and lactation. After weaning, exposure was conducted again during the mating and gestation period of the F 1 b litter production. Exposure was stopped a day or two before delivery
	0 Mating Interval	Mating continued for 15 days.
	0 Mating Ratio	Two females to one male
	0 Culling	Litters of more than 10 pups were reduced to that number by random sacrifice on the fourth day of the lactation period.
	0 Weaning	Litters were weaned 21 days postpartum
	◇ Production of Flb Generation Pups	After weaning the Flitters of pups, the dams were mated with different males selected from an ongoing chronic study.

- 0 Sacrifice, Gross Pathology and Organ Weights Males used to produce the F 1 a generation were sacrificed after mating and necropsied. Organ weights were recorded for brain, gonads, heart, kidneys, liver, lungs and spleen. A list of 3 1 tissues were removed and fixed for sectioning and staining including testes, seminal vesicles, prostate and pituitary.
- 0 Variations from current OECD 4 15 Protocol Guideline Significant variations include:
- . Number of pregnant females was about 10 rather than about 20.
 - Females were treated from the beginning of the mating period rather than from two weeks prior to mating in the F1 a mating.
 - . Only one dose group utilized.

Results

- Result Treatment did not significantly affect any measured reproductive parameter. There was a tendency toward an overall reduction in the number of pups born and weaned in both the F 1 a and F 1 b litters of the dosed group but this was not statistically significant.
- . NOAEL Authors conclusions:
0 FO (P) Generation 125 ppm
0 Fla Generation 125 ppm
0 F 1 b Generation 125 ppm
- . LOAEL Not stated in the report, apparently > 125 ppm

Remarks Field for Results ◇ Mortality During the Fla Mating and Gestation No mortality observed

0 Parental Data Fla Litter

Indices for Parental Parameters, F1a Litter (percent)					
Dose	Female Mating	Female Fecundity	Male Fertility	Female Fertility	Parturition
0	57.9	54.5	100	60.0	100.0
125	64.7	54.5	80	60.0	100.0
Number pregnant: control=6/10 125 ppm = 6/10					

0 Parental Body Weights Dams Fla

	Mean weights of dams on day indicated						
Dose (ppm)	G-15	Part	PP-1	PP-4	PP-12	PP-21	PP-28
0	409	366	361	359	379	373	348
125	377	348	343	346	354	359	339
G-15 = gestation day 15, Part= day of parturition, PP=postpartum day							

0 Progeny Data Fla Litter

	Number of pups that were			
Dose (ppm)	Delivered	Still-born	Cannibalized	Viable
0	65 (10.8) ^a	2	0	63 (10.5)
125ppm	43 (7.2)	1	0	42 (7.0)
a = mean per litter in parentheses				

0 Fla data

Fla		Survival Indices*		# Pups Weaned		21-day mean Body Weights (g)	
Dose	Live Birth Index	4-Day	21-Day	Male	Female	Male	Female
0	96.9	93.7	98.1	23	28	49	46
125	97.7	97.6	97.4	23	14	49	50

0 Fla Exam

All pups delivered appeared outwardly normal. An examination of each pup for external abnormalities was conducted at weaning and all pups were judged free of abnormalities.

0 Clinical Signs

None noted

0 Organ weights P-males

Males used to produce the Fla litter showed no difference in absolute or relative organ weights for brain, gonads, heart, kidneys, liver, or spleen. Absolute lung weights were reduced by 8 percent in treated males and the relative lung weights were reduced by 11 percent (statistically significant $p<.05$).

0 Histo-pathology

Microscopic examination of tissues from the males used to produce the Fla litter was unremarkable. The organ weight changes in lungs did not have a histopathological correlate. No adverse effects on reproductive tissues were noted.

- ① Mortality
During the Flb Mating and Gestation No mortality observed.

◇ F1b Litter Data, Parental Data	Indices for Parental Parameters F1b Litter (percent)					
	Dose (ppm)	Female Mating	Female Fecundity	Male Fertility	Female Fertility	Parturition
	0	92.9	69.2	100	90	100
	125	50.0	60.0	80	60	100
Number pregnant: control = 9/10, 125 ppm = 6/10						

- ① Dam Body Weights Body weights of dams were unaffected by prior treatment with dioxolane during the mating, parturition and lactation phase of the F1b litter production

◇ F1b Litter Data, Pup Data	Number of pups that were: [total (mean per dam)]			
	Dose (ppm)	Delivered	Still-born	Cannibalized
	0	91(10.1)	8(0.9)	0(0.0)
	125	64(10.7)	2(0.3)	0(0.0)

◇ F1b Litter Data, Pup Data	Survival Indices*		# Pups Weaned		21-day mean Body Weights (g)	
	Dose (ppm)	Live Birth Index	4-Day	21-Day	Male	Female
	0	91.2	91.6	67.7	29	13
	125	96.9	91.9	81.1	20	23

- ◇ F1b Pup Exam All pups delivered appeared outwardly normal with the exception of a control female pup that had **microphthalmia** of the right eye. An examination of each pup for external abnormalities was conducted at weaning and all pups were judged free of abnormalities.

- ① Clinical Signs None noted

- 0 Other The report contains no indication that the pups or dams were necropsied at sacrifice, it is assumed they were sacrificed without necropsy.
- 0 Comment Although treatment did not significantly affect any measured reproductive parameter, there was a tendency toward an overall reduction in the number of pups born and weaned in both the Fla and Flb litters of the dosed group. The lack of statistical significance may partly be a function of the low numbers of animals employed in the study. The possibility that the 125 ppm exposure represents a LOAEL for reproduction in the F 1 a and Flb litters cannot be excluded.

Conclusions

Remarks Field

Treatment of male and female albino rats with Dioxolane by inhalation at 125 ppm did not significantly affect any measured reproductive parameter. The tendency toward an overall reduction in the number of pups born and weaned in both the Fla and F 1 b litters of the dosed group suggests that the statistical significance could have been reached with a larger (n). Although 125 ppm was considered to be NOAEL by the authors of the study, the possibility **cannot** be excluded that 125 ppm represents a LOAEL for reproduction under these conditions.

Data Quality

- Reliability Klimisch Code 2. Study design, conduct and reporting are considered reliable to address the test endpoint although not conducted in accord with glp standards.

References

This study report was split into two documents by the testing laboratory. As it was a unified experiment with the same dams, both were combined for preparing this robust summary. The two reports are listed below.

- . Single Generation Reproduction Study with the Vapors of Dioxolane in Albino Rats Results of the FO and F 1a Generations, Status Report. IBT No. 663-05562 Industrial Bio-Test Laboratories, Inc. Submitted to PPG Industries March 1, 1976.
- Single Generation Reproduction Study with the Vapors of Dioxolane in Albino Rats Results of the FO and Flb Generations, Status Report. IBT No. 663-05562 Industrial Bio-Test Laboratories, Inc. Submitted to PPG Industries April 13, 1976.

Other

These results are supported by two oral studies from the same laboratory in four reports (reports were split by litter).

References for
supporting studies

1. Industrial BIO-TEST Laboratories, Inc.; One-Generation Reproduction Study With Dioxolane in Drinking Water in Albino Rats, Results of the FO Generation. (1975), EPA Document No. 878213523, Fiche No. OTS0205848.
2. Industrial BIO-TEST Laboratories, Inc.; One-Generation Reproduction Study With Dioxolane in Drinking Water in Albino Rats, Results of the FO Generation and Proven Breeder Males. (1975), EPA Document No. 878213524, Fiche No. OTS0205848
3. Industrial BIO-TEST Laboratories, Inc.; One-Generation Reproduction Study With Dioxolane in Drinking Water in Albino Rats, Results of the FO Generation. (1975), EPA Document No. 878213525, Fiche No. OTS0205848
4. Industrial BIO-TEST Laboratories, Inc.; Single Generation Reproduction Study With the Vapors of Dioxolane in Albino Rats, Results of the FO and Fla Generations, Status Report. (1976), EPA Document No. 878213527, Fiche No. OTS0205848

Definitions

- o Female Mating Index = $100(\text{number copulations}^i/\text{number estrus cycles required})$
 - o Female Fecundity Index = $100(\text{number pregnancies}/\text{number copulations})$
 - o Male Fertility Index = $100(\text{number sires}/\text{number males mated})$
 - o Female Fertility Index = $100(\text{number of pregnancies}/\text{number of females mated})$
 - o Parturition Index = $100(\text{number of parturitions}/\text{number pregnancies})$
 - o Survival Index, 4-day = $100(\text{number pups viable at lactation day 4}/\text{number of viable pups born})$
 - o Survival Index, 2 1-day = $100(\text{number pups viable at lactation day 2 l}/\text{number of viable pups at lactation day 4})$
- i) Only one copulation counted per estrus cycle.

Developmental Toxicology

Type	Developmental Toxicology
Test Substance	1,3-Dioxolane CAS Number: 646-06-0 Hoechst Celanese Lot UN1 166 Purity verified by analysis
Method	
• Guideline	U.S. Environmental Protection Agency Toxic Substances Control Act Test Guidelines - 798.4900
• GLP	Yes
• Year	1991
• Species	Rat
• Strain	Charles River CrI:CD®BR VAF/Plus
• Route of administration	Oral gavage in corn oil
• Doses	0, 125, 250, 500 and 1000 mg/kg
• Sex	Female, pregnant
• Exposure Period	Days 6 to 15 of pregnancy
• Frequency of treatment	Daily
• Control Group	Corn Oil
• Duration of test	15 days

- Statistical Methods

Maternal and fetal incidence data were analyzed using the Variance Test for Homogeneity of the Binomial **Distribution(1)**.

Maternal body weights, body weight changes, gravid uterine weights and feed consumption values, and litter averages for fetal body weights, percent male fetuses, percent resorbed conceptuses per litter, fetal ossification sites and percent fetal alterations were analyzed using Bartlett's Test of Homogeneity of **Variances(2)** and the Analysis of **Variance(3)**, when appropriate (i.e., Bartlett's Test was not significant ($P>0.05$)). If the Analysis of Variance was significant ($P<0.05$), **Dunnett's Test(4)** was used to identify the statistical significance of the individual groups. If the Analysis of Variance was not appropriate [i.e., Bartlett's Test was significant ($P<0.05$)], the Kruskal-Wallis **Test(5)** was used, when less than or equal to 75% ties were present; when more than 75% ties were present, Fisher's Exact Test(6) was used. In cases in which the Kruskal-Wallis Test was statistically significant ($P<0.05$), Dunn's Method of Multiple **Comparisons(7)** was used to identify the statistical significance of the individual groups. All other Caesarean-sectioning data were evaluated using the procedures previously described for the Kruskal-Wallis **Test(5)**.

1. Snedecor, G.W. and **Cochran**, W.G.. Variance test for homogeneity of the binomial distribution. Statistical Methods, 6th Edition, Iowa State University Press, Ames, pp. **240-241**.(1967).
2. Sokal, R.R. and Rohlf, F.J. Bartlett's test of homogeneity of variances. Biometry, W.H. Freeman and Co., San Francisco, pp. **370-371**(1969).
3. Snedecor, G.W. and **Cochran**, W.G. Analysis of Variance. Statistical Methods, 6th Edition, Iowa State University Press, Ames, pp. **258-275**(1967).
4. Dunnett, C.W. A multiple comparison procedure for comparing several treatments with a control. J. Amer. Stat. Assoc. 50:1096-1 **129**(1955).
5. Sokal, R.R. and Rohlf, F.J. Kruskal-Wallis Test. Biometry, W.H. Freeman and Co., San Francisco, pp. 388-389 (1969).
6. Siegel, S. Nonparametric Statistics for the Behavioral Sciences, McGraw-Hill, New York, pp. **96-104**(1956).
7. Dunn, O.J. Multiple comparisons using rank sums. Technometrics **6**(3):241-252(1964).

Remarks Field for 0 Age at Study Initiation
Test Conditions

- Females were approximately 93 days old at first dosing
- Males were approximately 24 weeks old when bred to the virgin females

0 Number of animals per
group

- 25 mated presumed-pregnant females per dose group

0 Test Substance
Preparation and
Analysis

- Test substance was prepared weekly at 4 concentrations (prepared twice) was analyzed for content of test material and each batch was analyzed seven days later to establish stability.

0 Analysis and Stability

Target Conc. (mg/gm)	Day-1 Conc (mg/gm)		7-Day Conc (mg/gm)	
	Prep 1	Prep 2	Prep 1	Prep 2
25	26	28	19	27
50	52	56	48	56
100	100	114	94	112
200	198	223	190	219

0 Vehicle

- Corn oil

0 Clinical Observation Performed and Frequency

- Viability observed at least twice daily.
- Additional observations for clinical signs of test substance effects, abortions, premature deliveries and deaths were also made immediately prior to intubation (days 6 through 15 of presumed gestation) and approximately one hour post-dosage. These additional observations were made once daily during the post-dosage period (days 16 through 20 of presumed gestation).
- The body weights of the rats were recorded at least once weekly prior to mating. Body weights and feed consumption values were recorded on day 0 and on days 6 through 20 of presumed gestations.

0 Mating Procedures

Following the acclimation period (approximately two weeks), 160 healthy virgin female rats were placed in cohabitation with 160 breeder male rats (one male rat per female rat). The remaining 20 virgin female rats were placed in cohabitation with breeder male rats that had mated during the first night of the cohabitation period. Female rats with spermatozoa observed in a vaginal lavage or a copulatory plug observed *in situ* were considered to be at day 0 of presumed gestation and assigned to individual housing. Female rats mated by the same male rat were assigned to different dosage groups.

0 Maternal Parameters Assessed During Study	Body weight, feed consumption and clinical signs during life. A gross necropsy was conducted on each rat. The intact gravid uterus was excised and weighed. The number and placement of implantation sites, early and late resorptions, live and dead fetuses, and the number of corpora lutea in each ovary were recorded. Uteri from rats that appeared nonpregnant were stained with ammonium sulfide to confirm pregnancy status
0 Fetal Parameters Assessed During Study	Litter size, placental weight, gross malformations, fetal body weight, sex ratio, body cross sections, skeletal examination. Live fetuses were sacrificed by immersion in the appropriate fixative. Approximately one-half of the fetuses in each litter were examined for soft tissue alterations by using a variation of Wilson's sectioning technique. The remaining fetuses in each litter were cleared, stained with alizarin red S(7) and examined for skeletal alterations.
0 Organs Examined at Necropsy	List not provided, stated in protocol that visceral organs that are abnormal will be noted and retained
0 Rationale for Dose Selection	The doses were selected based on a 14-day gavage study in non-pregnant rats of the same strain.

Results

- NOAEL& LOEL for Maternal Toxicity

0	NOAEL = 250 mg/kg/day
0	LOEL = 500 mg/kg/day Body weight gain
- NOAEL & LOEL for Developmental Toxicity

0	NOAEL = 500 mg/kg/day
0	LOEL = 1000 mg/kg/day Reduced fetal body weights and gross external, soft tissue and skeletal malformations or variations occurred in the 1000 mg/kg/day dosage group fetuses.
- Actual Doses Received

Not calculated, but very close to the target doses of 0, 125,250, 1500 and 1000 mg/kg/day
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- Maternal data

No rats died during the conduct of this study. No clinical or necropsy observations were caused by dosages of the test substance as high as 1000 **mg/kg/day**. The 500 and 1000 **mg/kg/day** dosages of Dioxolane caused weight loss in dams on days 6 to 7 of gestation. This initial weight loss was significant (**P<0.05**) for the 1000 **mg/kg/day** dosage group, and average maternal body weight gain was reduced for this group for the entire dosage period (calculated as days 6 to 16 of gestation), as compared with the control group value. Dosages of 250 and 500 **mg/kg/day** slightly reduced maternal feed consumption and the 1000 **mg/kg/day** dosage of the test substance produced a more marked reduction in absolute (g/day) and slightly reduced relative (g/kg/day) maternal feed consumption values during the dosage period as compared with the control group values. Significantly decreased (**P<0.05**) absolute feed consumption values occurred for the 500 and 1000 **mg/kg/day** dosage groups on days 6 to 7 of gestation. Administration of the 1000 **mg/kg/day** dosage of Dioxolane significantly decreased (**P<0.05**) absolute maternal feed consumption values for the entire dosage period.

- Fetal data

 - 0 Cesarean Data

Litter size was comparable in all groups. No Caesarean-sectioning parameters were affected by administration of 1,3-Dioxolane to the dams at dosages as high as 1000 **mg/kg/day**. The averages for corpora lutea, implantation sites, litter sizes, live fetuses and resorptions, and the numbers of dams with any resorptions were comparable for the five dosage groups. No litters consisted of only resorbed conceptuses

 - 0 External and visceral effects

The 1000 **mg/kg/day** dosage of the test substance resulted in significant increases in the litter and fetal **incidences** of externally evident tail malformations and vertebral malformations interrelated with the tail malformations and septal defects in the heart. The gross external and **soft** tissue malformations that were considered effects of the 1000 **mg/kg/day** dosage of the test substance were eight fetuses (from seven litters) in the 1000 **mg/kg/day** dosage group litters that had malformed tails. These malformations differed in severity and included constrictions of the tail, thread-like tail, short tail and absent tail; thread-like tail occurred for four fetuses (from four different litters) in this dosage group. Soft tissue examination revealed three fetuses (from three litters) at 1000 **mg/kg/day** dosage group litters that had ventricular septal defects as their only alteration. One **high-dose** fetus was found with a cleft palate.

- 0 Major skeletal defects
- Skeletal examination of fetuses from the 1000 **mg/kg/day** group revealed increases in vertebral malformations associated with the tail malformations (five fetuses ^{††} from four litters ^{††} had absent caudal vertebrae) and other alterations in vertebral and rib ossification that were probably interrelated with the more severe effects evident in the fetuses with visible tail malformations. These alterations included “scrambling” or reduced ossification of centra in the thoracic vertebrae (asymmetric, bifid, unilateral or absent ossification, fused **centrum** and arch), the lumbar vertebrae (fused **centrum** and arch, centra that were bifid or not ossified, and arches that were small or not ossified), absent sacral and caudal vertebrae and rib-vertebral malformations. These vertebral malformations and delays in ossification occurred in 4, 0, 0, 11 and 36^{††} fetuses from 3, 0, 0, 7 and 16 ^{††} litters in the 0, 125250,500, and 1000 **mg/kg/day** dosage groups, respectively. Of these alterations, significant increases ($P<0.01$) in the litter and fetal **incidences** occurred for thoracic vertebrae with bifid, unilaterally ossified or not ossified centra, lumbar vertebra with a bifid **centrum**, and absent caudal vertebrae. In the 1000 **mg/kg/day** group, the litter average for the average number of ossified metacarpal bones per fetus was significantly decreased ($P<0.01$) for this dosage group, a delay in ossification that is expected because of the significantly decreased ($P<0.01$) fetal body weight that also occurred for this group.

0 Fetal and Litter Effects

	Dose Group				
Effect	0	125	250	500	1000
Litters evaluated	25	24	25	23	24
Fetuses evaluated	405	388	382	362	376
Litters with altered fetuses	9	10	5	8	21 ^{††}
Fetuses with any alteration	17	10	6	16	52 ^{††}
Percent fetuses with any alteration per litter	4.43	2.44	1.52	4.16	13.78 ^{††}
Tail malformations, fetuses	0	0	0	0	8 ^{††}
Tail malformations, litters	0	0	0	0	7 ^{††}
Heart, ventricular septal defect	0	0	0	0	3 ^{††}
Vertebral scrambling, fetuses	4	0	0	11	36 ^{††}
Vertebral scrambling, litters	3	0	0	7	16 ^{††}
Vertebral/rib malform., fetuses	0	0	0	0	2
Vertebral/rib malform litters	0	0	0	0	2
Sternebrae alterations, fetuses	7	5	2	0	5
Sternebrae alterations, litters	4	5	2	0	4

- 0 Ossification Statistically significant reductions in the average number of ossified **metacarpals** per fetus occurred for the 500 and 1000 **mg/kg/day** dosage group litters, as compared with the control group values. This observation was considered biologically important and an effect of the test substance for only the 1000 **mg/kg/day** dosage group because: 1) no other statistically significant dosage-dependent delays in ossification occurred for the 500 **mg/kg/day** dosage group; 2) the values for the 500 and 1000 **mg/kg/day** dosage groups did not demonstrate expected differences reflecting the magnitude of the difference in the dosages provided (the 500 and 1000 **mg/kg/day** dosage group litters had averages of 3.12 ± 0.18 [†] and 3.10 ± 0.21 ^{††} ossified metacarpal sites per fetus, values that are not biologically different); and 3) the statistical significance of the value for the 500 **mg/kg/day** dosage group reflected the relatively small standard deviation that occurred for this group, as compared with the higher values for the standard deviations of the other dosage groups.

Remarks Field for Food Consumption
Results

Food Consumption per dam (gm/day)					
	Dose Level (mg/kg/day)				
Time	0	125	250	500	1000
Days 6-7	22.2	21.7	19.2	18.6 [†]	18.2 [†]
Days 6-16	22.4	23.6	21.3	21.4	20.7 [†]
Days 6-20	23.8	25.0	23.1	23.4	23.0

[†] = $p < 0.05$

	Dose Group (mg/kg/day)				
Parameter	0	125	250	500	1000
Pregnancies	25/25	25/25	25/25	23/25	24/25
Corpora lutea (per litter)	19.6	21.2	19.8	20.5	21.5
Implantations (per litter)	17.0	17.2	16.7	16.8	17.1
Litter size	16.2	16.2	15.3	15.7	15.7
Live fetuses	405	388	382	362	376
Dead fetuses	0	0	0	0	0
Early resorptions (total)	19	23	36	24	31
Late resorptions (total)	0	1	0	0	3
Resorptions per litter	4.4	5.8	8.6	6.5	8.3
Males %	47.8	52.0	50.0	52.1	49.5
Body weight (mean grams)	3.38	3.54	3.46	3.48	3.16 ^{††}

Conclusions

Remarks Field

Based on the results of this study, Dioxolane is not considered a specific developmental toxin. The developmental NOEL was found to be 500 **mg/kg/day** while the maternal NOEL was found to be 250 **mg/kg/day**. Neither deaths nor adverse clinical signs were observed in any of the animals. No adverse effects are anticipated for the conceptus in the absence of maternal toxicity.

The maternally-toxic 1000 **mg/kg/day** dosage of the test substance significantly reduced (**P<0.05** to **P<0.01**) fetal body weights for male and female fetuses and for the combined sexes. This dosage of the test substance also significantly increased (**P<0.05** to **P<0.01**) the litter and fetal **incidences** of externally evident tail malformations, vertebral malformations interrelated with the tail malformations and septal defects in the heart, and significantly reduced (**P<0.01**) the litter average for the average number of ossified metacarpal bones per fetus, an expected delay in ossification related to the significantly decreased (**P<0.01**) fetal body weights.

Data Quality

- Reliability

Klimisch Code 1. Reliable without restriction, study meets GLP standards and/or most requirements

References

1,3-Dioxolane: Oral Developmental Toxicity Study In Crl:CDOBR VAF/Plus's Presumed Pregnant Rats, Argus Research Laboratories, Inc. 905 Sheehy Drive Horsham, Pennsylvania 19044 Argus report # 508-002, May 9, 1991, submitted to Hoechst Celanese Corporation.

Other

A subsequent study was published which supports this result. In this study Dioxolane was administered by gavage every other day from days 8-20 of gestation as an aqueous solution daily doses of **0.14, 0.58** or 1.15 g/kg/day. Dioxolane administration was not associated with increased embryo or fetus intrauterine death rates or congenital defects at any dose level. The mid (0.58 g/kg) and high dose (1.15 g/kg) were reported to be associated with dose-related delays in fetal development and the high dose showed clear maternal toxicity. The developmental and maternal NOEL were both judged to be 0.58 g/kg/day under these conditions of every other day dosing.

References for supporting studies

Sitarek K, Baranski B, Berlinska B. The effect of maternal exposure to dioxolane on prenatal and postnatal development in rats. **Pol J Occup Med Environ Health** **5:159-66** (1992).

† = $p \leq 0.05$, †† = $p \leq 0.01$